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1 Tuesday, 8 December 2020

(10.00 am)

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SIR BRIAN LANGSTAFF: (Audio not available) ... anyone else because of Covid restrictions, I'm sorry that we have to restrict members of the public at the moment from coming to watch you in this location. They are, however, watching remotely, and we will have somewhere between 150 and 200 people, I would expect, watching what you are saying, able to tune in, or, for that matter, tune out, any time during the day. So that's your audience, if you like, the public in whose name this Inquiry is brought.

So, Mary, one of the members of staff, will now ask you to take the oath and then Ms Richards will ask you questions.

PROFESSOR IAN MALCOLM HANN, sworn Questioned by MS RICHARDS

MS RICHARDS: Good morning, Professor Hann. Can you see and hear me?

- 20 A. Very well, thank you.
- Q. I'm just going to ask you to give us an overview of
 your career. You undertook various house officer
 posts, I think, from 1972 onwards, and in the 70s you
 worked at the Royal Infirmary in Liverpool,
 Alder Hey Hospital and Manchester; is that right?

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membership exams in pathology during that period, round about 1981.

I spent a couple of months, two or three months, with Dr Kernoff and Professor Tuddenham at the beginning of that rotation, and the rest of the time I spent one year in pure haematopoiesis research, and the rest of the time in bone marrow transplant and leukaemia, almost entirely in adults but there were a few children.

- Q. When you took up your post then in the Royal Hospital for Sick Children in Glasgow in January 1983, how much experience would you say you had had by that stage in the treatment of patients with bleeding disorders?
- A. Apart from the fact that I'd had a gap of about 20 months, when I was mainly dealing with adults, with -- and doing research into leukaemia, bone marrow transplants, et cetera, I'd spent roughly, I would say, six or seven years, with exposure throughout the vast majority of that time, in haemostasis centres.

But of course I also -- during that period, I had to spend a year in neonatal training and a few months in other areas of -- well, six to 12 months in other areas of paediatric training, general paediatric training.

Q. Had you spent any time in a Regional Transfusion

A. Royal Manchester Children's Hospital, yes.

Q. In any of those posts, did your work involve the care and treatment of patients with bleeding disorders?

- A. I was in five separate haemophilia centres on six
 separate occasions. In Manchester, Liverpool, Great
 Ormond Street, the Royal Free and Liverpool Royal
 Infirmary.
- 8 **Q.** Prior to your appointment to the Royal Free, which 1'll come on to, were those rotational posts that you were undertaking? Did you also work in other areas?
- A. I worked in many other areas, particularly childhood
 malignancy, blood diseases, leukaemia in particular,
 and also I was doing my general paediatric training
 for seven years, and also pre-registration house jobs
 with Sir David Weatherall in Liverpool in haematology.
- Q. So your first time at Great Ormond Street I think was
 17 1978 to 1980, and then you moved to the Royal Free in
 18 1980 and worked there until the end of 1982.
- 19 A. That's correct, yes.
- 20 **Q.** In terms of your work at the Royal Free, what did that 21 entail?
- A. Again, it was a rotational thing because I was the
 first person to fully train in paediatrics and in
 haematology. I had to carry on training in adult
 haematology in particular until I took my final

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- A. Oh, yes, sorry. Yes, I spent a year -- in fact, in those days we were mandated to spend that period of time in Brentwood, and it was Edgware in those days, which moved later on to Colindale in north-west and north-east London.
 - Q. Can you recall roughly when that was?
- A. It would be -- it would be some time in the late
 1970s. I can't remember exactly.
- Q. You remained in post at Yorkhill until around August
 of 1987, when you moved to Great Ormond Street. Is
 that correct?
- 13 A. That's correct.
- Q. Then you were consultant haematologist at Great Ormond
 Street, paediatric haematologist, from 1987 to 2005?
- 16 A. The end of 2005, yes.
- 17 Q. You then spent I think roughly three and a half years18 working for a pharmaceutical company, for Bayer?
- A. I worked for four years with Bayer, developing
 longer-acting Factor VIII concentrates and the
 recombinant products and developing the new oral
 anticoagulant.
- Q. Then in 2000 or thereabouts you took up a post with
 the Irish Blood Transfusion Service as a paediatric
 haematologist?

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A. No, it was somewhat later than that. I had a period of time when I was -- between 2006 and 2010 I was at Bayer -- or the end of 2009, and then we moved to Ireland. My wife is a consultant here. And about six to 12 months later I took up a post as an intermittent locum and then a job share with the Irish Blood Transfusion Service, during which I spent six months as the national director.

Q. Now, I'm going to turn now to ask you some questions about the centre or the service at Yorkhill and your experiences when you arrived there in 1983.

When you took up your post, what was your impression of the service which you were taking on responsibility for?

A. Basically I found myself in the eye of a storm with not much calm. Dr Willoughby I had great respect for. He was a real workaholic and tended, to an extent, to plough his own furrow, to be a lone practitioner. It was very seriously under-resourced. Much of -- in fact, all of the junior or trainee staff were put in place by himself through charity and there were many challenges. I had seven areas of responsibility, of which haemophilia was one, and there were deficiencies in many areas which needed to be corrected, some of which were immediate safety issues.

well, I should have realised but I didn't fully realise how difficult medical politics could be. I was never very good at it, to be honest. But I was faced with battles on all fronts.

The immediate problem was that we didn't always get cover on the senior registrar haematology rotation. The adult trainees who needed to spend a few months with us, there were gaps when the staffing level was very poor. We had one person who was appointed as a leukaemia research fellow but in fact provided the clinical service. There was a threat initially, quite rightly in my view, from the Leukaemia Research Fund, that those posts -- which were essentially training posts -- would be removed. And the rest of the service was provided by clinical assistants, mainly on charitable money.

There was a shortage of social work and some resistance to the introduction of social work input. And there was no haemophilia centre. There was a lack of follow-up in certain areas. For instance, there was no multidisciplinary approach at all to the management of brain tumours. There was no follow-up regularly for children with haemophilia. There was no follow-up for the late effects of treatment of chemotherapy and radiation. So all those things had

- Q. What were the immediate safety issues that required correction?
- A. The laboratory, basically, was failing. And one of
 the main reasons why Dr Willoughby, as he told me,
 left was because of industrial actions throughout
 Scotland in the laboratories whereby the medics there was an intention to sideline the medics.

Basically, the quality control was inadequate and was repeatedly failing. The equipment was extremely old fashioned and, even in my experience, I couldn't remember its use. There was no computerisation. There was no adequate blood count machinery. The neonatal coagulation testing was very poor or inadequate or completely inadequate, despite the fact that we had a world-famous foetal medicine unit on site. Even, for instance, in blood transfusion we didn't even have any Coombs (unclear), which are the basics of cross-matching.

So a great deal needed to be done in that area with regard to safety. And with regard to other areas there were concerns over whether we had adequate cover to a trainee staff level.

- Q. What changes did you introduce in relation tostaffing?
- 25 A. It was very difficult. I didn't realise when I --

to be sorted out and three new clinics set up within a few weeks or months.

I was supported with that but running that level of department with that number of problems was extremely difficult.

- Q. In terms of clinical staff, I think there came a point, in 1984, when Dr Gibson joined; is that correct?
- A. Yes. That was a great relief.
- Q. In terms of facilities, what can you recall about the efforts made to improve the facilities and any moves in relation to the facilities for the haemophilia service?
- A. Well, with regard to the haemophilia service, it was really personnel that initially I could do something about. There was no space within the hospital offered to me. I did try very hard to set up a -- we had no haemophilia centre, essentially, and we had to rely on multi-use areas in the day care centre, and thankfully I was able -- I was facilitated in setting up a clinic which I could run every week if necessary, and at periods that did happen.

Basically, with regard to facilities, from the very outset I started fund-raising. We were a bone marrow transplant centre. We were essentially

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a national bone marrow transplant centre with very, very limited resources. So with Dr Alan Burnett, later Professor Burnett at the Royal Infirmary, I had to fund raise with him and approach the Health Minister in Scotland to get designation and funding. I had to spend far too much time fund-raising, pushing over piles of pennies in pubs and such like and eventually we were able to expand the laboratory and expand the office and data collection services. But it still meant that the clinical -- and also, by the way, when I first came I did not have any real designated beds and the service previously had relied upon the general paediatricians bolstering Dr Willoughby's expertise in haematology, and I was rather taken aback to find that they were taking decisions about my patients and that I didn't have a designated facility with fully trained nurses in any areas.

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So I had to address all of those problems, but I wasn't able, initially, Dr Gibson was more successful later on I'm pleased to say in setting up a haemophilia centre, but the lack of facilities was pretty widespread throughout the United Kingdom at the time.

When I went to Great Ormond Street in 1988 we

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cardiology, but other areas like infectious diseases and such like, hepatology, et cetera, were very lacking, and the consequence was that I had to rely on expertise from elsewhere.

- Q. One of the changes you instituted, which indeed Dr Pettigrew described to the Inquiry yesterday, was the establishment of some form of regular clinic system for patients with bleeding disorders. How often on average in 1983/1984 would a patient with severe haemophilia, whether haemophilia A or haemophilia B, have attended for a routine appointment once you had set up this system?
- A. Yes, routine clinic visits in haemophilia were not the absolute standard everywhere, but I was fortunate to have worked at the Royal Free where it was and, basically, I adopted a lot of their policies in inverted commas, such that if a patient was, say, a moderate haemophilia or a mild one I would expect to see them every year. They had very often not been seen for years previously.

With severe haemophilia it depended on the degree of severity and the degree of crippling, the degree of haemarthrosis that they had. I would expect to see them hopefully at least twice year, certainly every year, and for some of them it was required more

had a converted store cupboard basically as ourhaemophilia centre and nothing else.

- Q. You observe in your statement also about your time in Yorkhill that there were no specialists in the areas of infectious disease, hepatology, immunology or virology. What was the impact of that?
- 7 A. It was very difficult. I'd come from an ivory tower 8 into a situation where Scotland -- I had to say that 9 the Health Service in Scotland always worked better in 10 my experience than it did elsewhere, but one area that 11 was slow, and I experienced the same in Ireland I'm 12 afraid, is that developments were slow to come along. 13 One of the biggest gaps was the lack of sub-specialist 14 paediatrics. A lot of that was down to resistance 15 amongst the general paediatric consultant population.

Just taking immunology, for instance, I was not a fully trained immunologist. I'd been fortunate to be at the National Centre at Great Ormond Street. I had very good contacts there but I had to look after the patients with severe combined immune deficiency as well as everything else, also the patients with autoimmune diseases and so I had to rely a great deal on the contacts that I had there.

Within Yorkhill itself there were areas of great expertise in renal medicine, urology,

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often, say, every four months or so. During the period when we were having to improve communications, something I had to -- also had to work very hard on and didn't do very well with, then it was more frequent than that.

- Q. I'll come back --
- A. Sorry, also of course there was a completely open door policy with regard to the day care facility and I had great faith in Dr Pettigrew as an excellent communicator.
- 11 Q. I will come back later, Professor Hann, to ask you12 more about issues of communication.

In terms of Dr Willoughby's departure, other than the issue over industrial action to which you've already referred, what was your understanding of Dr Willoughby's reasons for moving and for leaving Yorkhill?

A. I think -- I only ever had a relatively brief conversation with him lasting perhaps 20 minutes or so. In that, he explained how despairing he was of the industrial or the threatened -- it never actually happened -- the threatened industrial action but also he was -- he had problems with a microbiologist consultant who had tried to pioneer rather pseudo scientific treatment of children with solid tumours

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using interferon, which was a flavour of the month for a period of time in the UK, and also he had struggled with resources and felt that he wasn't supported enough in regard to facilities and staffing and suchlike and had had to resort to the use of extremely generous charitable money, which I was quite surprised to find when I arrived and was just coming in when I had been asking for it. People in the West of Scotland were incredibly generous.

- Q. What were the out-of-hours arrangements from early 1983 onwards, particularly insofar as they would affect patients with bleeding disorders?
- A. Out-of-hours, the -- well, basically I was on call all the time. I had one week off during my first year there. It was supposed to be two weeks. I was called back after one week and there was no more cover from another hospital in that respect.

The patients who required -- most of the patients Dr Willoughby had very successfully started on home therapy and so we didn't see a great number of patients. It was really very infrequent for them to come up out-of-hours. They were seen on the ward usually and very occasionally in the casualty by the on-call person who would be one of the rotating senior registrars from the Royal Infirmary or Western General

properly resourced. Often one ran up against the great needs of other areas of paediatrics.

Q. You said in your statement that Scotland was slow to emerge in specialised paediatric care.

Can you elaborate upon that, particularly by reference to what you saw the position in Scotland to be at the time.

A. Yes. I mean, as I said, I think that the Scottish
Health Service worked incredibly well but it -- and it
is a relatively small country, very similar in size to
Ireland, and therefore you know if you work in a much
larger country there is much more drive towards
specialisation and regionalisation, if you like, as we
used to call it. Within Scotland itself, there was
a lot of resistance to the development -- I can give
you an example, paediatric oncology, really it took
years and years after I -- even after I left, for that
to be resolved adequately.

There was a tendency for surgeons to go their own way when managing solid tumour patients, often quite inappropriately, and really it's a question of how fast the Health Service responds to need, to developments and it was slow to develop -- to respond to developments like, for instance, bone marrow transplantation. Great Ormond Street itself didn't

Hospital or the Leukaemia Research Fellow on-call.

You know, it was very difficult to fulfil the on-call rotas adequately.

- Q. Do you know why the services at Yorkhill had been arranged or existed in such a way as to, as it were, lump in the care of patients with bleeding disorders with all paediatric cancer care?
- A. I don't know how that arose but it was pretty similar to what happened in other children's hospitals. It was part of the training, essential training, of course for senior registrars in haematology who were usually adult trainees, although some paediatricians came along later.

It was, along with solid tumours which at the time were also looked after by persons like me, although I had some, quite a lot of training in management of children with solid tumours, it took a long time to get a paediatric oncologist on board in Glasgow. There was a lot of resistance.

So you know there was -- these were, if you like, relatively new specialties. The specialty of paediatric oncology really didn't exist before 1971 but it took a long time with the institution of the UKCCSG but it took a long time thereafter for it to become, if you like, par for the course and to be

have anything approaching a proper bone marrow transplant unit until the early part of the 1980s despite the fact it had been going on for some years previously.

We were designated as a national centre in Scotland in around about 1983 -- sorry, 1985 but it took a lot of effort.

- Q. What did you understand the designation of the haemophilia service at Yorkhill to be within the broader UKHCDO context?
- A. Yes, I think, I don't know what the -- I've been asked this quite a lot and I really don't know the answer in a legal sense. The difficulty I think that arose, and personally I think it's a moot point because if you have children there and they need treatment I don't care what you are designated as, you provide the best available treatment. The fact is that Health Service managers in those days, and probably to this day, do not particularly like designation of specialist centres because specialist centres come with specialist pleading with regard to resources.

We were -- as far as my memory takes me, my contract says that I was going to be the director of the haemophilia centre. The patients needed that and we needed to improve the resources as did many other

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services at the time.

So to answer your question in a round about way, I did not regard ourselves as being a reference centre of the nature that the Royal Free was, which was in fact a designated international reference centre in that it had every possible resource. We were, thanks to Katharine Dormandy and people, we were very under-resourced but we were a haemophilia centre.

I did think that it was a bad idea that the UKHCDO split itself into various tiers and I actually made representations about that on a number of occasions because of the difficulties. You know, it was a whole different area with regard to communication and you relied on meetings and minutes of meetings (although doctors are terrible at minutes of meetings, I have to tell you, at producing them accurately) and, any question of what they actually meant, going back to them often was very difficult because you didn't have any idea what the discussions were.

In the end it became a doctors' organisation and in my view it should have been broader than that, but the fact was that we were a haemophilia centre, we were definitely not a reference centre in the Royal Free sense, but we were invited to some

period of time, which is why I needed Dr Gibson to come on board and why I had to spend so much time fighting politics and organisation and resource management and all the rest of it. It was -- I tried my best, but the biggest problem that I saw was with regard to communications. And I thought that I put in place adequate ways of dealing with that, because of my repeated absences from the scene, but clearly it was not adequate.

Q. Now, one of the first tasks you took on when you joined Yorkhill was to produce standard operating procedures or written protocols or policies.

As I understand it, you produced those across the range of your various different responsibilities; is that right?

A. Yes. It took a great deal of time. My family certainly didn't appreciate it.

There were no standard operating policies in the department other than the protocols that were produced for the treatment of leukaemia essentially. I needed to begin at step 1 from the very beginning and write those policies out. It took, in the end, between -- well, certainly up to six months to do that.

I think that they were very valuable. People

directors' meetings.

Q. You did attend, I think, the annual general meeting
 from October 1983 onwards, representing Yorkhill. The
 earlier minutes seem to suggest that Dr Willoughby did
 not attend. Do you know why? Was anything about that
 ever raised with you?

A. No, it wasn't, but I knew him quite well actually, and he attended a lot of meetings about leukaemia and bone marrow transplants and all that sort of thing. The fact was, he was a single-handed workaholic. Getting away to London actually improved a lot when we were allowed to use aeroplanes and newfangled things like that, but it was still -- most of these meetings were held in London or even more difficult places to get to. London became relatively easier.

But, to my knowledge, he attended such meetings very infrequently, if at all.

- Q. What proportion of your time was spent dealing with bleeding disorder patients as opposed to your multiple other responsibilities in the period from 1983 to 1985?
- A. Well, of course it varied enormously depending on the
 current crisis or acute, say, serious head injury or
 whatever it might be. It was rarely more than 10 or
 20 per cent of my time. And it was an inadequate

said so at the time, I don't know if they were being kind or not, but it was certainly essential that they were there, because there were people who were not getting adequate information themselves. Doctors, nurses, et cetera, at the time did not have access, in a time when there was no internet and email and all the rest of it, and when we relied a great deal on attendance at meetings, and very long delay in general publications. So, yes, I regarded it as a very high priority.

- 11 Q. It may be difficult for you to remember the detail,
 12 and we don't have the texts themselves, but can you
 13 recall what the policy or guidelines in relation to
 14 the care of patients with bleeding disorders -- can
 15 you recall broadly what it said or what its purpose
 16 was?
- A. Yes. I mean, basically it would be a preamble about the -- a little bit about the history of haemophilia, about how it presents, about the various levels, the various tests and what tests you perform and when. And all of that -- not exactly lifted from the Royal Free and Great Ormond Street but -- mainly the Royal Free -- certainly what was the standard mantra and practice there.

It would also include -- and I know that --

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I believe that others have explained -- the use of other products such as DDAVP, EACA, tranexamic acid, cryoprecipitate, et cetera, and it would have gone into quite extensive detail on the use of that because, of course, some of them were of real safety concerns when it came to treating children.

Then you would break it down into areas which were actually designated to quite a large extent in the book that I gave out to as many children that I saw as I could living with haemophilia, where it described what was serious, what might not be so serious and what sort of levels you need to achieve, et cetera.

- Q. Now, one of the main changes which you instituted in terms of treatment policy was to move away from the use of commercial concentrates and to use almost entirely and, in later years, I think entirely SNBTS concentrates. Can you just tell us a little bit more about your thinking?
- A. Yes, I had in fact discussed this briefly with Dr Willoughby, who, as you saw from my statement I hope, basically preferred that, the use of commercial concentrate, because of this purity, essentially, and to a limited extent its availability.

 I was impressed with the Blood Transfusion

short period of time and we changed over.

And I can give you the reasons why if you wish.

Q. Yes, please.

A. Basically, again, I've been asked lots of times if I've watch a television programme -- I'm almost sure I didn't -- that talked about the skid row blood donors et cetera in America. I just didn't have time during that period of much on call and young children et cetera. But I was very well aware, having worked in blood transfusion, of the mantra, which obviously to this very day I still respect as being of vital importance, which is that blood and blood products should be donated by people who are very effectively screened, which they were not in those days, very effectively tested, as far as you can, which they certainly are nowadays, and not in those days, and altruistic and with no inducement whatsoever.

There was no doubt we knew at the time that -- I need to put it in context a little bit, because there's obviously a lot of concern about non-A, non-B hepatitis. All of the concern or almost all of the concern in the early 1980s was of hepatitis B. We know to this day that hepatitis B can still be transmitted. We had a case in Ireland of a blood transfusion donor three years ago. It is still not

Service because very soon, within days or a few weeks I had contacted them and asked them what the score was, why are we using all this commercial concentrate -- I mean, it was ironic for me because we had been -- we, I mean the consultants that I worked for in the UK -- had been fighting for years for self-sufficiency, and on many, many occasions I had heard this both formally and informally, and I was told that -- I contacted I think it was Dr Crawford, who was our main liaison, and he was extremely helpful and he said, you know, "If anything, we will give you priority. I can't guarantee you absolute full supply but I think we can do."

So right from the word go, and within days or weeks at the most, I stopped all use of Factor VIII concentrate from abroad.

There was, at some stage, and I've been asked about this a number of times, some residual commercial concentrate. And my memory, which may not be correct, is that we sent this for destruction. Which is quite a difficult decision in one way because it is costly stuff and all that, but that paled into insignificance as far as I was concerned.

So the answer is that the people on home therapy used up their supply within a relatively

a solved problem. It certainly wasn't a solved problem then.

So, you know, we knew that the type of donors that were being used in America were very high risk in this context. I could not contemplate continuing the use of commercial concentrate having, ironically, heard for years how much the people in England wished to do so. So yes, I knew -- I felt that it was incredibly important. It was with some dismay that I learned later on that the Health Life questionnaires in Scotland and elsewhere allowed prisoners to donate, another population that we know may not possibly sometimes be altruistic and certainly do have a higher risk of a virally transmitted infection.

- Q. You mentioned in your answer a few minutes ago, in relation to your conversation with Dr Crawford, that he said effectively he could accord you a degree of priority or preferential treatment. Was that because yours was a paediatric service?
- A. I think so. I mean, I've always tried not to do down
 adult services but one -- certainly at Great Ormond
 Street it was commonly the case that Blood Transfusion
 Services said: yes, we'll give you priority. You
 know, that's an ethical/moral issue which is difficult
 to come to terms with sometimes.

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I think that from my vague memory of the conversations with Bob Crawford at the time that he was very keen that we, if you like, came on board and that the Blood Transfusion Service, which was very well run by John Cash and others, that they should be self-sufficient and they should be that very soon.

I can talk about self-sufficiency if you want because it's used far too loosely.

Q. I might come back to that at a later stage if I may, Professor Hann. I want to focus on Yorkhill specifically for the time being.

Your changes in relation to treatment policy as well as switching to SNBTS from commercial concentrates included, I think your statement says, a policy to use factor concentrates conservatively and only when necessary; is that correct?

A. Yes. The difficulty in very early 1983, I told you about my knowledge of HIV -- HIV or AIDS before that. Basically, we did not know about dosage effects, about infectivity and suchlike. So I and others, I discussed it with people at the Royal Free and elsewhere, felt that it was a reasonable idea to be more conservative with the use of concentrates, even in those on home treatment.

In that respect, you'll know from my statement

infrequently or not previously treated patients or PUPs or previously very low volume treated, infrequently treated patients.

So, yes, there was a change in policy where it was feasible and I am very happy to talk about cryo if it's appropriate to talk about the use of cryoprecipitate at some stage.

- Q. Yes, my next question, in fact, is going to be what use was made under your new treatment regime from 1983 onwards of cryoprecipitate and for which categories of patient?
- A. Basically I think what you may well have heard before I haven't been able to watch all the evidence. We used it in patients with von Willebrand's disease where it was necessary and DDAVP wasn't appropriate, type 3, type 2B or serious bleeds or where we couldn't get levels down, which was a real problem during this period.

We used it in patients with mild haemophilia if the bleed was not serious and we used it in new patients as they came along. It was discussed with patients who were receiving concentrates and there were some who wished to change to cryoprecipitate and my memory is very vague on this. I think there were just one or two, maybe even three, people, families,

that I actually terminated, after discussion with the families, the prophylaxis in some patients.

In retrospect, I very much doubt that that made much difference but the fact was that I could see -- I was an initial sceptic. Dr Willoughby was before his time. He took on board the work that others should and I should have taken on board, from Sweden mainly, Dr Yeoman Imanu(?) and Marie Nilsson showing that prophylaxis could work but in the patients that he was treating it wasn't working.

Dr Liesner very importantly showed later on that it could work if you persisted and if you gave adequate amounts and you achieved levels above 1 per cent. It wasn't working in those patients so at that time I didn't see any need to continue it.

I also wanted to be absolutely sure that I didn't beggar the nation's supply on the basis of a treatment that really wasn't working.

On the second aspect of this, to cut through the protocols that I wrote, I had very strongly re-emphasised the need to consider the use of alternatives such as DDAVP, et cetera, and for a period of time there was a consensus, I believe, in the people that I spoke to anyway, to use cryoprecipitate for newly diagnosed or very

who wished to revert to the use cryoprecipitate and as far as I know that did happen. But the use of cryoprecipitate in children is very limited and I can expand on that if you wish.

- Q. Well, I think your statement addresses it to some extent but what are your particular concerns about the use of cryoprecipitate in children?
- A. I'm afraid it's going to be a bit of a long answer but I'll try and keep it as short as I can.

I think that there was some very important work from Scotland by Charles Forbes in the period when we were switching over to concentrates, when I was a trainee around about 1976, which showed that there was no tendency in reduction in premature death during the cryoprecipitate period, which basically started very soon after I went to medical school in 1966. I think it was talked about in about 1964, something like that.

So during that period the commonest cause of death in haemophilia patients was intracranial haemorrhage and, as you probably know, one of the commonest causes or needs for treatment of haemophilia patients in childhood is head injuries. Bearing in mind that initially we did not have access to CT scans, et cetera, and therefore we had to rely on

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history, signs and symptoms, children who have head injuries, as everyone knows, often are a bit sleepy afterwards. They might vomit. How are you supposed to address that? You address it by giving a treatment which is well tolerated and where you can rely on the dosage being what it actually says on the bottle, which is not the case for cryoprecipitate.

There are problems, obviously, I can talk about home therapy separately if you want because that's a whole separate issue with regard to cryoprecipitate but, basically, cryo was a large volume, it took a lot of time to draw up, it disenfranchised the families who had had their lives revolutionised in the mid -- around about 1976 period.

On top of that, it wasn't a safe treatment. The efficacy was extremely difficult to manage during this period. We had lots of laboratory problems, as did everywhere, with overreaction to the HIV era. It was extremely difficult to get reliable, fast Factor VIII levels, for instance, and so you needed to give cryoprecipitate following a head injury or an internal bleed or severe haemarthrosis, probably every 12 hours and sometimes every eight hours.

It took time to do this, usually up to an hour or more. The volume could not be just passed in, it

A. Correct.

- Q. In relation to those families who, to the best of your recollection, asked to revert to cryoprecipitate, you did use cryoprecipitate or you did accede to their request?
- A. I wouldn't have refused their request no, that's for sure. But, you know, they were aware, many of them. I mean, one of the problems I had and my problems were nothing compared with theirs was that with the -- a lot of these patients had been coming for a number of years. Some of them had already received cryoprecipitate and knew what it was like and so the knowledge to a certain extent was there but not entirely and so it was necessary to make them -- I wouldn't scare them to death and say there's a risk of death with giving this because that risk was very small, but not insignificant. But it would be explained to them what the downsides were. Most of them were already well aware of that.
- Q. In terms of the commercial concentrates, do you know what the arrangement had been prior to your arrival in January 1983 to procure commercial concentrates and were there commercial arrangements that you had to terminate?
- A. It worked, as far as I remember, like this. There

would cause transfusion overload in children.

Finally, round about one third to a half of patients had side effects. These could be very, very frightening for children. They consist of rigours and shakes and feeling dreadful and, occasionally, these could be life-threatening. I had experiences with Dr John Martin in Alder Hey with a patient who developed severe anaphylaxis requiring chlorpheniramine, aminophylline, steroids and adrenaline, and that patient would not have -- would almost certainly not have survived at home. So, you know, with regard to home therapy I can come back to that if you wish. It was, in my view, not an option. I had not, in all those times in haemophilia centres, including (unclear), seen anyone who had -- in a young child, had been able to institute home therapy.

So it was a useful treatment, it did help quite a lot, especially for minor bleeding, more minor bleeding disorders, but it was extremely difficult to use in children.

Q. But you did use cryoprecipitate for children with von Willebrand's, you did use it for children with mild or moderate haemophilia if DDAVP was not an option -- is that correct -- because you tried to avoid concentrates?

were very, very few occasions on which I needed to do this and my memory is vague, because the only situation in which it would occur was there was one patient who developed an inhibitor and had a massive requirement, and there were several other patients who needed fairly urgent synovectomies done which required a great deal of replacement. So there may have been a few occasions, and very few occasions, on which we reordered commercial concentrate and that would be through the senior chief in the lab in contact with the pharmacy.

I didn't -- it wasn't an era of resource management. I didn't hold budgets or anything like that, and I was never restricted in the use of anything because of financial costs or of that nature.

- Q. The evidence we've seen so far tends to suggest that Yorkhill didn't have a system of what other clinicians have referred to as batch dedication, and that that was something that appears to be considered perhaps for the first time in late '84 and early '85. I can take you to some minutes if necessary that refer to that?
- A. Yes, no, you are absolutely right. It was later on and, again, I was having very regular -- I can't remember them all or even a small proportion of them.

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I was having very regular discussions with Mark Crawford who was an excellent liaison and at some stage or another he did say we -- John Cash and him had discussed whether we could dedicate batches to particular patients; in other words, you know, when they came up for their therapy, whether it be at home or in the hospital, usually at home, they would be allocated the batch that was for them.

I think it was quite a bit later on. It may have been in '84, probably was in '84 in fact, that that occurred and I think that a small number -- in a small number of patients that's what we received.

- Q. In terms of the changes that you instituted to the home therapy programme they were no longer doing it on a prophylactic basis, how was that change introduced? Was every patient or family brought into the centre or was it the next time they attended to collect their supplies?
- A. Yes. I mean, these are patients who continued to have many bleeds, sometimes even on a weekly basis or whatever. They would be seen in -- I had to rely on Dr Pettigrew and I relied on her a great deal.
 I probably relied on her too much. But the discussions took place mainly between herself,
 Sister Murphy and the family. If the family had said

been very frequent.

- Q. Professor Ludlam also told us about meetings that would take place with the Scottish Home and Health Department. Are those meetings that you participated in as far as you can recall?
- A. It's possible but personally I doubt it because I was the representative for paediatric haematology and oncology on these -- I think they called them -- special advisory groups of what was then called the Home and Health Department. What with everything and that, I doubt that I would have attended this as well, but I tried my best to keep up through minutes but also, more importantly for me, via Gordon Lowe, who was a really very helpful, always very helpful, contact at the Glasgow Royal Infirmary.
- Q. If we go to the second page, please, Soumik, and could we look at the paragraph in the middle of the page, so (ii). Just a little bit further down, please.

So we can see here Dr Cash has introduced a paper, various matters are being discussed, and then at (ii) it says this:

"Members discussed the suggestion that the production of cryoprecipitate could now be reduced. Dr Ludlam said that cryoprecipitate was preferred in the treatment of children at present, because of the

we wish to continue, we would have done so. But I didn't really see much point, to be honest.

I didn't want to run into a situation where we ran out of concentrate as a consequence. But that wasn't the main reason.

Q. I want to look at one document with you,Professor Hann.

It is PRSE0001556, please, Soumik.

- A. Will I see that?
- 10 Q. Yes, it should come up on the screen in a few moments.11 Can you see that, professor?
- 12 A. Yes, I can.

Q. So if we just zoom in to -- thank you, Soumik.

So we can see it's the minutes of a meeting of directors of the Scottish National Blood Transfusion Service and Haemophilia Directors on 2 February 1984. We can see that you were in attendance.

Before we look at the substance of the minutes, Professor Ludlam last week told us that one of the features of local arrangements were regular meetings between haemophilia directors and SNBTS. Is that an arrangement that you participated in? We can see you here but did you participate on a regular basis in these meetings?

A. I certainly went to some of them but it wouldn't have

new danger of AIDS. Dr Hann concurred. A policy seemed to be emerging however to use less cryo for haemophilia A patients. It was agreed that a certain minimal amount of cryo was required and Dr Cash pointed out that TDs could produce it in emergencies."

I don't know whether you have any recollection of this meeting or discussions at the time, professor, but you will see there the suggestion appears to be being made, which you agree with, that cryoprecipitate is, by February '84 at least, regarded as the preferential treatment for children because of the risk of AIDS. Do you have any observations on that?

- A. My first observation is that it's pretty typical of medical minutes which you can interpret various ways. I presume that what is meant here is that it was preferred for newly diagnosed patients with haemophilia. It wasn't the preferred method for all children. And obviously for those other groups that we've discussed it was appropriate. And later on, when heat treatment became available in, I think for us, early 1985, it was regarded, by myself anyway at least, as cryoprecipitate being less safe.
 - Q. So do we correctly understand your evidence to be that you don't agree with what's said here if it's interpreted broadly as applying to the treatment of

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children generally? 2 A. I don't agree with two things. First of all, I don't 3 remember ever saying we wanted to use less cryo. And 4 secondly, I don't agree that it should be used for all 5 children, no. Q. You have mentioned heat-treated products. Can you 6

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- recall what, if any, involvement you had in trials of heat-treated products in advance of them becoming more widely available at the end of 1984?
- A. I don't recall being involved in any such trials, although of course we were crying out for what we called, and still do, pathogen reduction techniques. It takes an epidemic/pandemic crisis for these things to happen, unfortunately. We've been talking about it for years. We wanted it, we needed it. But I was concerned.

I think I contacted Dr Cash at the time that before taking on any heat treatment in children, as with any other pharmaceutical or blood product, that I needed information from the adult sector first. For two reasons: first of all, I needed to be sure that it was well tolerated and it wasn't going to cause even worse reactions than we occasionally saw with concentrate, and; secondly, I was very concerned, as was everybody, that neoantigens, in particular

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1 purpose of it was?

> A. Yes, I don't really recall. I can only speculate that it was looking for neoantigens and particular Factor VIII inhibitors as well as the other tests that are indicated there, which are looking for transfusion transmitted infections. But also it would almost certainly have included what is called in the jargon "recovery"; in other words, looking at whether the levels of Factor VIII that you achieve in the patients are those you would expect from the dose that's written on the bottle.

Q. You go on to say:

"I do still wonder whether, in the 'virgin' cases the frequency of testing will be enough to document the non-A, non-B problem especially as LFTs, EBV, CMV, Hep B, Hep A etc are not mentioned. In addition, I wonder how useful a vague efficacy study will be."

Can you recall what your concerns were in those regards?

A. Yes, I suppose as far as the scientific data collecting aspect of this is concerned, it was like many of the haemophilia studies at the time, and subsequently, based on what is just a collection of anecdotes if you like, some of which I've already

inhibitors to Factor VIII, did not develop.

It usually is the case that such occurrences occur very quickly and thus I was able, as far as I remember, to be able to take on the heat treatment product in children quickly. But, you know, developing inhibitor is a disaster in haemophilia. It's a life-long, potentially, issue. It requires treatment which, at that time, was not available, largely anyway, as far as curative treatment is concerned, and therefore I was very sceptical, but at the same time very hopeful, if that doesn't sound stupid.

We'll just look at your exchange of correspondence with Dr Cash on this topic.

Soumik, it's PRSE0003840, please.

This is a letter dated 19 December 1984 from you to Dr Cash. You thank him for a letter. The heading is "SNBTS [heat-treated] Factor VIII". You

"We'll do our best to take samples from our children although I cannot guarantee their frequency will be great. Certainly this will produce a significant increase in workload for us."

Can you recall what the request had been to you in relation to the taking of samples and what the

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mentioned.

As far as the other tests there are concerned, I was still obviously concerned as to whether the heat treatment process would be sufficient for other viruses than HIV. I was not, to the best of my knowledge, and I have racked my brain about this, provided with what I subsequently was involved with with Bayer, which is so-called spiking studies, in other words, looking at how well this actually worked in practice with regard to reduction of virus, of various viruses. And I think that I just had to rely on word of mouth and, thankfully, subsequent experience that HIV was labile in this short heat treatment process.

I wasn't fully aware at the time, although I know subsequently, that it wasn't adequate for hepatitis C but -- well, what was subsequently known as hepatitis C, but again, I wasn't provided with that information, specific scientific information.

Q. You say in the second paragraph:

"I would just like, at this time, to express my own ethical and professional doubts at the way in which this major change has been instituted. The impression generated to the parents is that their children will now be protected from AIDS and that

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prior products were dangerous."

I understand, I think, professor, the first part of what you are saying there. You were not, as I understand your evidence, entirely confident that the SNBTS heat-treated Factor VIII would provide protection from AIDS, but --

A. That correct.

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- Q. -- what was your concern about parents getting the impression that prior products were dangerous? Because wasn't that right? They had been, at least for some recipients.
 - A. Yes, I've not put it very well there. I mean, basically what I'm trying to say is that it should be put in context and that whereas SNBTS products were relatively safer than commercial concentrates as far as I was concerned, unfortunately, due to failures of Health Life questionnaires in particular that the --I don't know actually when the breakthroughs occurred in Scotland in retrospect in relation to this timing of this letter but I'd always -- I'd never used the word absolutely safe with regard to any pharmaceutical or blood product.

There's a reason why the ABPI bans the use of the word "safe" full stop in all of its guarantee publications from drug companies, et cetera. There is

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through proper development processes and to use it with consent, if the parents agreed.

I don't recall ever giving such product but would have considered this if there were good data available with regard to efficacy, neoantigens and transmissibility.

Q. You can take that down, thank you, Soumik.

In terms of previously untreated patients, was there any particular protocol or policy that you introduced from 1983 onwards to address how decisions should be taken and what treatment should or should not be given to a previously untreated patient?

A. Yes. Obviously this was following what I call the legacy period and at that stage I made sure that I at some stage spoke to the families of new patients There were very few of them. I don't know how many but it would be a very small handful during the period of time I was there by chance, and because of the population that we were dealing with was of the order of 3 million and probably half a million children.

Basically, I would speak to the -- and with Anna usually and Sister Murphy quite often, and sometimes the social worker if available, and explain to them about haemophilia and all the background of the severities and the treatments and the risks,

no such thing as a fully safe product, but I did believe, as you know, that there was a relatively good degree of safety, relatively speaking, with regard to the SNBTS product.

Q. I just want to ask you about one further document relating to heat-treated products.

> Soumik, it's WITN4183002, please. Go to the next page.

We can see here there's a reference, it's a letter from Dr Pettigrew to Armour, which refers to having been sent a supply of heat-treated Factor VIII, although it appears from the letter that it wasn't in fact used. We also saw with Dr Pettigrew yesterday a letter in which she and you were named on a clinical trial exemption entry in relation to Armour's heat-treated product, again in March of 1984.

Do you have any recollection of why either you were approaching Armour or Armour was approaching you in relation to heat-treated trials at this stage? A. I don't have any specific memory but we were, all haemophilia directors probably, were in regular touch with the commercial companies of various sorts. I didn't have any reason to favour Armour over anyone else. Basically, we were pretty desperate at this stage to get heat-treated product that had been

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et cetera, and would say that in the first instance we would be using cryoprecipitate, and give them Peter Jones' excellent book Living with Haemophilia, The Haemophilia Society leaflets that were available and put them -- and give them the contacts for The Haemophilia Society and encourage them strongly to join and, if possible, to attend their meetings and, if possible, to attend the World Federation of Haemophilia, which was almost unique at the time in allowing or encouraging the patients to attend.

Q. You've emphasised in your statement the importance of training and advice for clinical staff in relation to the risks of transfusion-transmitted infections.

Had there been any training programmes, as far as you're aware, prior to your arrival in 1983 at

A. I don't think there were many training programmes. It was sort of on-the-job training, if you like, which unfortunately happened a fair bit in that era, in many areas.

One thing that's important to say probably is that because of a disaster that occurred with hepatitis B causing deaths, several deaths in a laboratory facility, a policy of -- an excellent policy, which still exists today and sometimes gets

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Yorkhill?

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forgotten, was the so-called universal precautions with regard to samples and how they should be treated and getting rid of this business of high-risk, et cetera, which I always did not like and did not think it was suitable. Unfortunately, people resurrected it again with Covid-19. Some people never learn.

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Basically, in addition to that there would be, obviously in the adult area, there would be training with regard to sexual transmission, et cetera, but that was as -- and we were transferring the patients once they were giving self-therapy and in secondary education it did not arise. But the risk of transmission of infections for those giving treatments and dealing with blood spills, et cetera, was emphasised throughout and Sister Murphy was extremely good at doing that. I was present when she was doing that on a number of occasions.

Q. Dr Pettigrew told us yesterday that she was not aware of even the idea of non-A, non-B hepatitis, at least in those terms, until she attended a scientific meeting in 1984 at the Royal Free. I'm going to ask you in a little while separately about non-A, non-B hepatitis but what if any efforts were made from 1983 onwards to provide junior medical staff with access to

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essential that that happened. There were very little funding for that. We needed people to attend the British Society of Haematology, the World Federation of Haemophilia which, unfortunately, was in places which cost a lot to get to, et cetera, like Brazil, and also, for me, the seminal meeting, which was the American Society of Haematology, and in that respect I basically had to contact the various -- all of the companies producing concentrates or anything else in relation to haemophilia and ask them for funding on a rotational basis.

Also, there were occasions where we discussed haemophilia on the ward rounds and at the -- and, rarely, at the hospital meetings, which were unfortunately often overwhelmed with leukaemia and solid tumours. I'm sure that they have better training nowadays but I was just one person.

MS RICHARDS: I'm going to move on to another topic, sir, and I note the time, so should we perhaps take a break now?

SIR BRIAN LANGSTAFF: Yes, we take a break mid-morning. Now, it may be unlikely that you want to talk to anyone about the evidence you have given or are yet likely to give but you mustn't. You can talk about anything else you like.

journals or to ensure that they were kept up to date with medical and scientific developments?

A. It's absolutely true to say that the level of training that I found when I came there was not adequate and the methods of training consisted largely of, as I said, on-the-job training. You know, if a problem occurred it was addressed and there might be further presentations within the hospital, et cetera, of such things.

What I did was probably not adequate but it consisted of -- there was very, very little reading of journals, something which I was just not used to, and having been an avid reader myself and in that era there wasn't much -- many other ways of getting information. I ordered again, this was back to charity again, I ordered a series of journals, about ten in all, and I began marking them up with a page stuck to the front of it and circulating them around the department's doctors. I would take home a massive pile to read at the weekend and then mark those up and send them around and sometimes we would photocopy seminal articles and give them to the trainee staff.

In addition to that, I had noticed that Dr Pettigrew in particular had not been facilitated in attending international/national meetings, and it was

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1 A. I'll talk to the cat. He's outside. 2 SIR BRIAN LANGSTAFF: We'll take a break for half-an-hour. 3 So let's come back say 11.45, shall we. 4 MS RICHARDS: Thank you, sir. 5 (11.18 am) 6 7 (11.46 am)

(A short break)

MS RICHARDS: Professor Hann, I wanted to ask you a little more now about non-A, non-B hepatitis. Your evidence in your statement is to the effect that non-A, non-B was regarded as no more than a minor illness at the time you started at Yorkhill.

Now, it's clear from the evidence you've given so far, that hepatitis B was regarded a very serious condition, indeed. What was the basis for your belief or understanding or assumption that non-A, non-B hepatitis was minor?

A. Right, I didn't put it very well, then, to be honest, with you. What had happened in the latter part of 1979 and 1980 was that hepatitis -- non-A, non-B hepatitis had come much more into the forefront of publications, et cetera, but unfortunately, and it brings up a problem that I think we struggled with and I, you know, believe that I failed to a certain extent to communicate adequately, and that is the level of

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uncertainty associated with certain problems.

The point about it is that we knew by this stage, and I'm sure you've had evidence of this, that probably your first dose of concentrate was enough to cause -- to give you a dose of hepatitis. And latterly we knew that was hepatitis C, from 1989 onwards.

What that did not tell you, and what led to all the uncertainty, was what the prevalence of persistent

What that did not tell you, and what led to all the uncertainty, was what the prevalence of persistent infection and liver damage was. Subsequently, we knew that it was eradicated endogenously by the patient themselves in between 25 and 40 per cent of cases. And to give you an idea of the uncertainty of what clinical problems would ensue, during my time at Great Ormond Street between '88 and 2006, during which time some of the patients would be in their late 20s and being followed up jointly, just one patient required treatment for hepatitis C hepatopathy.

We also knew at a later stage, probably in 1985, from the studies of Pierre Mannucci and Lou Aledort in over 100 patients receiving liver biopsies, that the frequency of liver diseases in concentrate patients was no greater than in patients treated principally with cryo. And at this time, they therefore said, and this was the mantra if you like:

instance occurred around about 1968, that it fell to a very low level by 1990, when hepatitis C was actually found. We know, and it was my experience, that very few children had symptomatic episodes of hepatitis with non-A, non-B. So, you know, both diagnosis, prevalence, persistence of infection were very uncertain.

Then, you know, we were faced with the two papers from Sheffield, the second of which was the most worrying and turned out to be -- although it was a perfectly proper publication, and I knew Sir John Lilleyman very well, we worked together for years in London, it turned out to be very misleading. I think it was five children, two of whom were only two and five years of age, who bravely received liver biopsies and who were shown to have significant persistent liver disease.

We knew subsequently, of course, that the progression was 1 per cent per year and that, as I said, spontaneous clearance could occur frequently in non-HIV patients and that cancer was less than 1 per cent per year, and life expectancy subsequently shown to be reduced by only between two and six years.

So there were a great deal of uncertainty, which -- it was our job, if you like, to transmit

there's no indication to alter current therapy because of concern over transfusion-transmitted infection from plasma production.

That finding, clinically anyway, was reviewed again in 2003 by Mannucci, who confirmed those findings.

Basically, the problems we had were of uncertainty. There was no diagnosis. The mantra was if you found that the liver transaminases function tests were more than doubled for six months, then you could think that that might be non-A, non-B hepatitis. But, of course, that turned out to be a completely insensitive test and many patients were affected, despite not having those findings. I don't actually remember any patient who would have fulfilled that criteria. So we couldn't say you had non-A, non-B hepatitis even.

It was probably multifactorial, and Purcell -- and I was trained to a certain extent by

Dame Sheila Sherlock, who was the guru of hepatology and viral hepatology, and basically they said: don't use the term "hepatitis anything" because we don't know what it is, it's probably multifactorial, it could be due to a ragbag or multiple viruses.

We knew subsequently that the peaking of

information to patients without causing very unnecessary anxiety. I don't mean that in a patronising or patriarchal way. It's damaging for families to be told very uncertain information and to be hung the Sword of Damocles above them for many years. And many years would be the case. The average age in the Sheffield adult cohort was around about 40 or something like that.

So it was information that I hoped and assumed, incorrectly, had been transmitted largely over the years, sometimes more than a decade prior to my coming there. I did not spend enough time, having made that assumption, reinforcing information with the legacy patients, if you like. And despite the fact that product information or patient information leaflets were present in the packaging from the late 1970s, Haemophilia Society leaflets became available, World Federation Haemophilia meetings were sponsored, and patients attended, very unusually, Peter Jones' book subsequently, within the next year or so, dealt with it, and all the other things, like parent support groups, independent support groups, and a new clinic, all of these were not adequate to adequately provide the information, which I expect you wanted to me to talk about. And we did not -- I did not. I made

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assumption that knowledge was better than it was, that information had been provided that had not been adequately provided. And I wasn't proactive enough in seeking information as opposed to doing what we were taught to do, which is just ask open questions.

I'm sorry, that's a very long-winded answer but a very important question, for me.

Q. Professor, thank you, and that addresses to some extent the questions I have been intending to ask you about the question of what information was and wasn't provided to patients at Yorkhill.

Can I just take you back to the question of what your understanding was in the period 1983 to 1985 in particular in relation to non-A, non-B hepatitis.

I understand what you say about uncertainty, and you've referred to various subsequent studies, and what may have been learnt from them. We have seen, for example, materials from Dr Kernoff from 1979 and 1980, I think -- certainly from 1979 -- characterising non-A, non-B hepatitis as being a serious disease with long-term consequences.

Would you disagree with or would you have disagreed with that characterisation in the early 1980s?

A. To a certain extent I would disagree because we didn't

about this patient with leukaemia", or whatever, "The potassium is 7 and I don't know why", and so on. Am I to go in and tell the parents that this is something that is incompatible with life? Or do I explain to them, as I tried to, that we would sort this out as fast as possible? And in that circumstance, you can sort it out as fast as possible, but in this context of non-A, non-B hepatitis, et cetera, you could not.

We basically -- let's say, haemolysed blood sample is common in children with difficult blood samples. That's just one example of many where you could put your anxiety -- dump it on a family and expect them to cope with it without full and long explanation, in a circumstance where you already know what the likely causes are and you can be, to an extent, reassuring. And where you have some appropriate therapy, which did not, of course -- was not, of course, the case with hepatitis C as it -- on non-A, non-B hepatitis in most cases.

So my answer in the same is incomplete, and I hope that explains it a bit better.

Q. So would this be a fair understanding of your evidence, and please correct me if this is not fair or not correct, in your view, in the early '80s it would know -- if you'd added the words "could be" or something like that, I would entirely agree with it. What we didn't know -- and I'm not being evasive, really -- we did not know the prevalence. You know, how do you communicate that? The answer is: inadequately. In my case. And I regret that.

The fact is that -- can I give you just two very brief examples of the problems that one faces as a paediatrician? And I'm not asking anyone to feel sorry for me.

The fact is that, you know, when it came to new variant CJD, what were we supposed -- this is much later obviously -- what were we supposed to tell people? We had no diagnostic test. We didn't know who could get it. We knew, eventually, that you could transmit it. We knew that similar diseases, like Kuru could take 30 years to develop. What were we supposed to do? Tell people that, "Maybe over the next 30 to 40 years you may get a debilitating life-threatening horrific neurological disease"? The fact is that these are damaging psychological bits of information.

And just as another quick example, it would be common in those days to, say, go to the ward -- and I can just give you one instance of many, where you go to the ward, the registrar says, "I'm a bit worried

have been correct to say that non-A, non-B hepatitis
 could be a serious disease with long-term
 consequences?

A. Yes.

5 Q. It could --

A. You know, how does one then define, you know, when it becomes a serious illness and therefore you are hanging the Sword of Damocles over somebody because for sure I can tell you now I am not going to be the doctor that does liver biopsies on children with haemophilia in that era. It's a dangerous procedure.
It leads to deaths.

Q. Can I leave aside for a moment the question, you are right it logically follows, as to what you might tell patients and just think about what was understood, albeit with the uncertainties you have identified, you did understand or you would have understood that it could lead to chronic hepatitis and could lead to liver disease but the prevalence of that was one of the areas of uncertainty.

Is that a correct summary? I know it's only a summary of your evidence.

- **A.** Yes, I agree. The prevalence and the definition were uncertain. The diagnosis of it.
- Q. You have referred to the Sheffield study in relation

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- to children. You were, I think from your earlier
 evidence, probably also aware of Professor Preston's
 Sheffield work.
 A. Yes.
- Q. Were you aware of what I think we've seen described in minutes of the UKHCDO's Hepatitis Working Party of the
 Sheffield Royal Free collaborative liver biopsy study
 that was ongoing I think certainly in 1980?
- 9 A. I wasn't aware of it. I know that Peter Kernoff told
 10 me I think or at least I found out somewhere, I think
 11 he told me, that they had had one death related to
 12 a liver biopsy which I wasn't aware of that study, but
 13 at least I can't remember being aware of it, sorry.
- Q. If we move then from the question of what your understanding was and your awareness of uncertainties to the question of what should have been said to patients, again as I understand your --
- SIR BRIAN LANGSTAFF: Just before you go there, is one way
 of summing it up as to your knowledge after 1978
 before 1983/84 that although you didn't know how
 prevalent it was, you knew there was a real risk that
 blood products might transfer hepatitis non-A, non-B?
- 23 A. Yes, we did know that.
- 24 **SIR BRIAN LANGSTAFF**: And you knew there was a real risk that it might be serious?

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- questions", and I did not spend enough time going
 through non-A, non-B hepatitis with people who had
 been around for years and attended on many occasions.
 That's my fault.
 - Q. In terms of Dr Jones' book, we can look at the texts if need be but they are what they are, they say what they say, the first book, which would have been the one available certainly in 1983, had been published in 1974 and deals only with Australia antigen hepatitis B.
- 11 A. That's correct.

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Q. The second edition was published some time in 1984 and certainly does make some reference to non-A, non-B hepatitis and, indeed, to AIDS.

Do you happen to know when in 1984 it was published?

- A. I don't but it should be easy enough to find that out from the ISBN.
- 19 Q. I'm sure it will be, thank you.
- A. I do remember ordering it as soon as it was available
 because I thought it was really good and it probably
 made up for some of my own inadequacies.
- Q. Would you accept as a matter of principle that whilst
 there may be lots of other valuable resources that you
 can provide to patients, ultimately it's the

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1 **A.** Yes, but, you know, I'm sorry to keep rambling on about uncertainty --

- SIR BRIAN LANGSTAFF: I'm just looking for a simple way of expressing what you are saying and --
- A. I have a slight caveat in that I'm not in any way
 criticising the Sheffield Children's study which was
 completely valid, but it was extremely misleading.
 Yes, I was aware that it could be serious, yes.
 - MS RICHARDS: If one then comes on to the question which you addressed a few minutes ago, of what was said to, and what should have been said to children, to families at Yorkhill, as I understand your evidence, you are acknowledging that you didn't make sufficient proactive enquiries to establish what they had been told; is that right?
- 16 A. Correct.
- 17 **Q.** And is this right: although information was available
 18 to patients in the form of Dr Jones' book and the
 19 various other sources of information that you referred
 20 to, there wasn't any proactive or systematic approach
 21 to ensuring that your patients received that
 22 information from the clinicians at Yorkhill?
- A. Not proactively, no, other than the fact that
 I immediately recognised the need for a review clinic
 at which they were asked, "Do you have any other

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- obligation of the clinician to spell out to patients
 the risks and benefits of treatment rather than make
 assumptions that they might read materials elsewhere?
 - **A.** Yes, absolutely and the buck stopped with me not with Dr Pettigrew.
 - Q. I want to ask you then about your knowledge of AIDS and, in particular, to ask you about a symposium you attended in June 1982 because you're the first of the oral witnesses that we've had I think to make reference to that. This was something called the Second International Symposium on Infections in the Immunocompromised Host held in Stirling in June 1982.

What was the nature of the event and at whom was it, as far as you are concerned, predominantly aimed?

A. This was an incredibly valuable meeting which was
 initially started off by the European organisation for
 research into the treatment of cancer.

In that era, and it was my particular area of expertise and specialisation, deaths from the treatment as well as the disease itself but particularly the treatment of leukaemia were far too high and infections in the, as they called it, the immunocompromised host, in other words, usually cancer patients, leukaemia patients, et cetera, of all ages

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was a very important topic and a matter of governance which I had in particular took on board for, initially for Manchester but more widely thereafter. It was my particular area of expertise.

So I attended it really for that reason and I won't easily forget it because I managed to become very ill during it with my carditis and non-A, non-B hepatitis for the second time due to coxsackievirus and I only say that because my memory of it is very clear despite the fact it was very long ago.

It was a very, very shocked meeting, indeed, and it's impossible to exaggerate that. There was one thought in particular and in those days it wasn't called AIDS or HIV or anything else. It was called GRID, which I think gives you the idea of the emphasis in those days: gay-related immunodeficiency disease. And, to cut a very long story short, amongst this extremely frightening presentation and others, there was a list of persons who were affected, and at the very bottom of that list were either two or three persons with haemophilia who -- I don't even know if it got into the final publication but I remember it vividly.

So that was my first indication that there was a problem. Of course, one didn't know what the

cities in the US. It is affecting homosexual men, intravenous drug users of either sex, and Haitian refugees."

Then if we go on please, Soumik, to page 22, we look at the last paragraph on this page we can see there reference to the mortality rate:

"Acquired immunodeficiency disease has high mortality rate. 13 of 42 patients in our series have already died. Nationwide, half the patients have died."

Then there's a description of what is said to be the typical course of the illness.

Then if we go on two pages please, Soumik, we can see if we look into the first main paragraph about six lines down the absence of treatment is referred to at that stage:

"Treatment of infections in these patients is hampered by the severity and apparent irreversibility of immunodeficiency", and further details are given in relation to that.

Then if we go to the next paragraph please, Soumik:

"The aetiology of this acquired immunodeficiency disease is not known", and then there's discussion about cytomegalovirus and other

co-morbidities were in those people but there were lots and lots of theories as to the cause of it and most people felt that it was multifactorial at the time.

Q. I'm going to invite you to look at one of the papers from it, not least because we haven't looked at it in any of the hearings so far.

Soumik, it's PRSE0002220, please.

This is the subsequent publication of materials relating to the symposium which you attended in June 1982. We can see the date there being 1983.

A. Yes.

Q. If we could turn please, Soumik, to page 18 I think it will be of what you have, we can see there in fact this is entitled:

"AIDS, acquired immunodeficiency syndrome, infection and neoplasia in homosexual men and intravenous drug addicts."

We can see the authors there and I think you referred in your evidence to the Penrose Inquiry to some of those authors.

22 A. Yes.

Q. It opens with:

"We're experiencing an alarming epidemic of an acquired immunodeficiency syndrome, AIDS, in certain

hypotheses. Then at the bottom of the page this:

"A role for an infectious agent is suggested by the fact that the disease occurs in people in whom infections are readily spread. Blood or body secretions would appear to be [if we go to the next page, top of the next page] potential vehicles of infection but since many people are exposed and only a few people develop clinical symptoms factors such as differences in host resistance and inoculum size may be important."

Then it goes on to discuss the particular position of patients with enlarged lymph glands. Then if we look at the very final paragraph on that page, Soumik, we can see there are a discussion of the devastation of the illness:

"Those who take care of these patients realise how devastating this illness is. The early events need to be identified by prospective studies of high-risk groups", and so on.

So that's the written paper. You described in your evidence to Penrose, I think, about this being a bombshell essentially in terms of the nature and severity of this new disease; is that right?

24 A. Yes, it was, yes.

Q. Again, you told the Penrose Inquiry, we can look at

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the transcripts if you wish, but I hope these are accurate summaries, that you left the conference thinking that it was likely that this new disease was caused by a new viral agent and that it might possibly be relevant to patients with haemophilia; is that fair?

A. Correct.

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Q. Now, you said again in your statement and evidence to the Penrose Inquiry that those who were in attendance at this conference were, in particular, those who were interested in leukaemia rather than those who were -who were clotters, as I think you put it.

Were there other haematologists present, as far as you can recall, who had the broader interest which included the care of people with bleeding disorders that you had?

A. No. Other haematologists, yes, but paediatric haematologist are more different, as I expect you would notice by now, that we had to wear several hats in those eras, whereas haemophilia treaters tended to be in the main or totally haemophilia treaters and they would not have attended this conference. But there were a large -- it was the best, by far, conference of its nature in the world, and it was attended by many microbiologists and infectious

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press, The Lancet, the New England Journal, et cetera, et cetera and -- I don't recall how many there were but there were many publications. It was a very, very hot topic from then onwards, and in many areas.

I mean, haemophilia treaters don't just read publications about haemophilia. I mean, they read the general haematology literature and they would be expected to keep up with the general literature such as the New England Journal and The Lancet.

- Q. You referred to there having been reference to two or three cases with haemophiliacs you remember from the conference, and we know that it was in only the following month, July 1982, that the CDC in its MMWR publication drew international attention to those cases.
- A. Yes.
- Q. Was the MMWR a publication that you had access to at the time?
- A. No, it wasn't. It's a daily reading for me now, in transfusion, but it certainly wasn't widely read in that era. It required -- it was like a newsletter almost. It still is. But it's very widely read throughout the world now. But in that era it certainly was not.
- Q. You were, at this time, in 1982, still at the

disease doctors. And there were, I think, even people from the SNBTS there who gave a talk. That can be checked but I'm pretty sure that's correct.

So it wouldn't have been something that haemophilia doctors would necessarily have been aware of, because of this meeting, but it was part of the burgeoning knowledge that began to explode at that time.

- Q. Is it something that should have been shared with haemophilia doctors? I don't mean necessarily by you, Professor Hann, but you have referred more generally in your statement, and indeed your evidence this morning, to issues about communication, information sharing, availability of information. How would the news from this conference, how could it have been disseminated more widely to those treating haemophiliacs?
- 18 A. It was quite difficult because it was -- it's not 19 exactly a niche area, by any means, but it was 20 produced -- in those days, it was quite common to 21 produce a book of a meeting. Unless you bought that 22 book or disseminated that book, you wouldn't know 23 about it. But I'm sure you've found that there were 24 a whole series of publications. I would expect 25 haemophilia doctors to be aware of the general medical

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1 Royal Free, although I don't know whether you were 2 still working with Dr Kernoff and Professor Tuddenham 3 at that stage?

- A. No, I was there, I'm almost certain, in the early part
 of 1980, and although it was a joined-up department,
 I wasn't working with him or Professor Tuddenham for
 the previous nearly two years.
 - Q. Do you recall any discussion at the Royal Free in the latter half of 1982 about this new illness, of AIDS?
- 10 A. Oh, yes, there was. And I can't remember any specific 11 details but there were a lot of -- the Royal Free was 12 very well set up for all sorts of things, including 13 lectures and education and seminars and all the rest 14 of it, to which all of the Haematology Department 15 would attend. And I can't remember exactly when but 16 certainly during that period there would have been 17 discussion and presentations on the subject.
 - Q. You've referred to the New England Journal of Medicine, and we know there was an editorial in the New England Journal of Medicine in January of 1983 by Jane Desforges that looked at AIDS and haemophilia.
- 22 A. Yes.
- Q. Is that something you think you would have read at thetime?

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25 A. Yes, I think so, although I don't know -- I did not

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- 1 know of her particularly as I don't think she was 2 a haemophilia doctor. But that doesn't negate the 3 findings. 4 Q. The Lancet. At the end of January 1983, again, there 5 was an article in The Lancet about AIDS and 6 haemophilia. Would The Lancet also have been on your
 - general reading list at the time? A. Definitely, yes.
- 9 Q. I want to pick up what was being discussed amongst 10 haemophilia clinicians in January 1983 about AIDS by reference to the minutes of a meeting that you weren't 11 12 at, professor, and we wouldn't have expected you to 13 have seen, but it's a useful summary, rather than 14 looking at lots of different documents, of a state of 15 knowledge.
- 16 A. Yes.

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17 Q. It's PRSE0002647, please, Soumik.

> It's a document I think you are aware of now. It's "Notes of meeting with Immuno at London Airport -24 January". It's January 1983.

If we go to the fourth page, please, Soumik, we can see from the list of attendees that at this meeting there were multiple leading Haemophilia Centre Directors, not just Reference Centre Directors but others as well.

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If we can just go to the top of the next page, we can see then there's reference to:

"... incubation period ... appears to be six months to two years.

"In the UK, so far only one or two cases have been reported ..."

And then there's a discussion about protocols in the US and reference then to the New England Journal of Medicine.

Now, I'll check with you, professor, but I have assumed, is this correct, that you weren't aware of this meeting at the time?

- A. Can you just tell me the date again, please?
- Q. Sorry, 24 January 1983. So just as you had moved to Yorkhill?
- A. No, I have subsequently been made aware of the meeting but I knew nothing about it at the time or its
 - Q. I have taken you to it, professor, just, as I say, a convenient summary of information that was being shared at least amongst a number of haemophilia clinicians at the time. Given what's set out here, which was, I think, not knowledge unique to those attending the meeting but had been fairly widely reported, would you agree with this, that certainly by

Soumik.

Q. Including, in terms of Scottish presence, Dr Ludlam. If we go to the previous page -- and we pick it up in the last two paragraphs on that page, please,

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Professor, this is in the context of a discussion at this meeting in London, with Dr Craske summarising the current position in relation to AIDS.

You'll see there that the minutes record or the notes record it being said that:

"Up to 10 December ... some 800 people had been reported as suffering from the AIDS, and there was a 45 per cent mortality.

"Ten haemophiliacs in the US have been affected and five have died. The youngest was aged 7. All cases have had prolonged treatment with Factor VIII, but there is no specific implication of one particular product or batch."

Then it goes on to refer to three cases involving blood and blood product transmission in non-bleeding disorder patients, so transfusion of platelets, one of which was a 20-month old child who developed a possible AIDS state. That was a case that had been reported in the MMWR in December of 1982, professor.

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1 the end of January 1983, by which time you have had 2 the New England Journal and The Lancet publications, 3 it should have been understood that the likeliest 4 cause of haemophiliacs being infected, although the 5 numbers were still relatively small, was the use of

6 blood products? 7

- A. There was a slight glitch in the sound there, sorry. 8 What date by did you say? 9
 - Q. By the end of January 1983.
- 10 A. Yes, I think, again, we don't -- didn't know what the 11 prevalence was likely to be but, yes, we would have 12 been aware by this stage that it was becoming -- it 13 had become clear that haemophilia persons were 14 affected.
 - Q. So haemophiliacs were at risk. You didn't know necessarily the extent of that risk. And at risk of being infected with something which was known to have a very high mortality rate?
- 19 A. That very high mortality rate is, of course, in people 20 who have other risk factors, so we didn't know what 21 the mortality rate was going to be but it was 22 frightening enough as it was, yes.
- 23 Q. I think -- again, appreciating you didn't know exactly 24 what was being said at this meeting, but we see it 25 being reported by Dr Craske that in five of the ten

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1 haemophiliac cases reported in the States the patient page, where they say: 2 had died. 2 "... any developments of significance will be 3 3 A. Yes. circulated to you before then ..." 4 Q. I want to then take you to March 1983, to a document 4 That's before the next, AGM, in October: 5 that you should have received at the time. It's 5 "... as soon as information is available." 6 6 a communication from Dr Craske. Do you have any recollection of receiving this 7 It's HCDO0000517 001, please, Soumik. 7 letter, Professor Hann? 8 8 So we can see this is a letter dated A. I don't have any specific recollection but, as I've 9 9 22 March 1983: said in my statement, I hope, that we were asked --10 10 "Dear Director ..." I don't know if you are asking whether we responded to this but there were a lot of requests for information 11 Our understanding is that it was sent to every 11 12 haemophilia centre director. 12 to be sent to centralised organisations and other 13 13 It refers to recent discussions in the organisations, et cetera; and all of us, Dr Pettigrew 14 Hepatitis Working Party and a meeting of Reference 14 and myself and others, were very concerned about 15 Centre Directors and the circulation of various 15 disseminating any information other than truly non --16 enclosed papers. The aim of which was to set up 16 I can't think of the word -- anonymised, sorry, a system for the reporting of possible cases of AIDS. 17 information. If anonymised information was requested, 17 18 18 If we look in the second paragraph, we can see from a proper organisation, then we would have 19 the criteria for reporting are set out. 19 complied with that. 20 And in the third paragraph, Dr Craske says: 20 Q. You were being asked here, as all directors were, to 21 "We ... strongly urge you to collaborate in 21 effectively report to Dr Craske any suspected cases 22 22 reporting cases of this syndrome ..." of AIDS. Do you recall whether you did so? 23 If we look at the next page, we can see it's 23 A. No, we didn't have any suspected cases of AIDS. 24 from Dr Craske, Dr Rizza and Professor Bloom. 24 Q. But would you agree that the reason you're being asked 25 I should perhaps draw attention to the top of the 25 to do this by UKHCDO at this time -- when I say you 73 74 1 1

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I mean all directors -- is because of the recognition that it was likely that AIDS was transmissible by blood products? That was the very reason for asking directors to look out for possible cases.

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A. Well, the main reason, yes, is the answer to that but the main reason is in fact to know about the prevalence, the natural history. You know, we needed to know how to manage these patients.

- Q. The thrust of your evidence to the Penrose Inquiry was that it was really in the second half of 1983 that it hit you and other directors that AIDS was going to be a significant issue. Why do you put it as the second half of 1983 rather than early in 1983?
- A. I think there's, from my point of view, there's been a little bit of conflation of HIV infection with AIDS which I think is important to be clear about.

We didn't know the natural history, as I've just said. We needed to know that in order to be able to inform people and to know how to manage, how to follow-up people or how to use prophylaxis in children and so on. So the answer is the big clue came in May of 1983 and I think you may have already heard about all that, Montagnier's description.

There was a fair bit of not scepticism but lack of understanding as to whether this was really the

agent at the time and so on. So as far as the virus was concerned, my own view and that of many others was that he'd cracked it and he got the Nobel Prize and all that but it wasn't really for another year until we were absolutely sure 100 per cent that that was causing the problem.

It was a transmissible disease. It was going into haemophilia persons who had been treated but we did not know -- it appeared that it was different in haemophilia. I expect you have had other people tell you all this so I'll be brief but it wasn't the same in haemophilia. We didn't have any patients who were wasting, we didn't have patients with the Kaposi's sarcoma. In fact, it was very uncommon in haemophilia because it was due -- I mean, AIDS is not caused by one virus. It was due to the HHV8 virus subsequently we knew, and so it was multifactorial. It was always thought that it was multifactorial and it turned out that it was.

So, I'm sorry, a long-winded answer. We knew that it was transmissible. We knew that HIV had entered sadly the haemophilia population. We did not know yet the natural history of the disease or its truth pathogenesis and we had no therapy.

Q. What was the point in time, doing the best that you

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can, professor, that you thought the parents of children with haemophilia had the right to know that the treatment that they were giving their child might transmit a fatal disease?

A. I prefer to use a different tense if possible: might have transmitted. There was little evidence subsequently. I mean, I'm not answering this very well. I wasn't aware, and I'm still not aware, as to whether any of the product that was used from 1983 onwards transmitted HIV. I think that my attitude at the time was that it was becoming more and more likely that HIV had been transmitted and that at some stage during 1983, especially after the publications and the finding of the virus, controversial though it was and continued to be, that discussions should begin at that time.

There was obviously common knowledge at this time, unfortunately reflected in the pariah status that many haemophilia families found themselves in, that HIV was a problem in the haemophilia population. It would have been unconscionable in a way that haemophilia families did not know that, certainly during the latter part of 1983 and, again, we discussed it as appropriate and passed on what information we could.

indeed in a Haemophilia Society Bulletin, which may or may not represent the risks in a way that a clinician thinks is accurate, it's for the clinician to explain their understanding, imperfect though it might be at a given time, to the patient?

A. Yes, although that sounds easier than it is, but you're absolutely right. I would entirely agree with you, which is why we set up parent groups and they set up their own groups and we had a social worker. It's a process in family communication. It's not just the almighty doctor coming in and pontificating on something. It's a process over a period of time and a development of trust and an ability for people to ask you questions when you don't know exactly what they know or what they want to know, no matter how open your questions are.

So yes, we did use The Haemophilia Society. We used their leaflets, we used their meetings, we encouraged people to go to the World Federation. I'm not going to run through it all again but it's not just -- and obviously the reason I set up the clinic was so that that facility was there and we didn't just rely on ad hoc visits.

Q. In the course of 1983 and 1984, how often would the clinics be clinics that you were taking and the

Q. Is this right, that it's not until after May 1983 that you would have thought it incumbent upon you to explain issues relating to risks of AIDS to families?

A. That's not quite true. I did -- if -- during 1983, I can't remember all the dates, I'm sorry, when it was in the meeting and all that. There was a welter of information, a lot of it very unhelpful, in the media. The families were extremely well aware of it and would ask questions about it.

But it's very tough sometimes to give information in the face of lots of lack of knowledge. That would be passed on as it was available and was very often in the media as soon as it was -- I mean Montagnier's findings were in the media before I knew it. It's not a question of keeping it from them or being lacking in transparency. I have nothing to -- we have nothing to hide. It was a question of how to provide adequate information. It wasn't simply giving them massive anxiety in a situation where they may well not have had it. That's the problem.

Q. Would you accept, I think, going back to a question I think and an answer you gave a few minutes ago, that patients -- it's not for patients to try and understand risks from the basis of what might be reported in the Mail on Sunday or The Observer or

1 conversations, conversations you were having as 2 distinct from Dr Pettigrew and Sister Murphy? 3 A. We mainly did it together. There was a shortage

A. We mainly did it together. There was a shortage of rooms apart from anything else. Dr Pettigrew and I -- Sister Murphy would quite often not be able to attend because she had other duties. We didn't have a dedicated nurse or a dedicated social worker or a dedicated psychologist. All of those were available and were quite good but intermittent to an extent.

To answer your question precisely, I can't but, the clinics were set up initially monthly but were frequently more often than that and I would attend them at least twice a month and sometimes on a weekly basis during the period when we were informing those families that had not had an opportunity to discuss it already in the day care centre with Dr Pettigrew.

- Q. Dr Pettigrew's evidence yesterday was that there was no policy as such about the provision of information in relation to AIDS to patients or their parents. Would you accept that?
- A. Well, there was no written policy but the policy is the policy of paediatrics and the -- it's an ethos rather than a policy, if you like. It's a question of an open door policy, first of all, first and foremost, transparency and honesty and to be as full

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We discussed it on numerous occasions. I didn't write down a policy because it depended on the family and their availability. It was a very difficult decision which we did not get -- and it's my fault -- we did not get entirely right with regard to the process as opposed to the policy, if you like, as to how to contact, how best to contact people, how best to see them, how best to get through to them all of the information that we had and was essential on all sorts of things like universal precautions, et cetera. So, no there wasn't a policy but it was a question of multiple discussions between us, which we agreed between us. But it was my responsibility.

- Q. It might be said that the thrust of Dr Pettigrew's evidence was that rather than proactively contacting patients to discuss risks of AIDS with them, the approach was reactive. If questions or concerns were expressed by parents, they were then addressed and there might then be a dialogue and conversation that would ensue. Is that your recollection?
- 22 A. It depends what you mean. Do you mean before or after 23 we got the results?
- 24 Q. I'm talking about the period before you got the results; so '83/84? 25

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stored. This was not absolute routine. As I said to you before, my interest was in infection in the immunocompromised patient so I'd done a lot of virology follow up of such patients, and I knew that to an extent where the samples were available and where they were adequate, which they often weren't for children because they were large samples, we knew that they were there.

We knew that there was a lot of development going on with regard to testing. All virology testing -- no virology testing is perfect. There is no such thing a virology test which is absolutely specific and accurate and sensitive, so there were real problems in the initial phase of development of the serology for HIV, with regard to false positives and false negatives, and it would have been incredibly unhelpful to pass on information in the face of that.

We still have problems in Blood Transfusion Services. For instance, if you go as a blood donor today you will be asked to sign a form that says, "I consent to my blood being tested for anything you like for the next 30 years or more". It's essential that we can do that for the sake of the donors and the recipients.

So the pre-counselling testing did not happen

A. The answer to your question is correct.

Q. That brings me then to the question about the results and the process by which you discovered that a number of patients, former and present, at Yorkhill tested positive for HTLV-III.

We heard yesterday from Dr Pettigrew her recollection that you received a letter from Dr Follett which set out results that hadn't positively been sought as far as she was aware by the haemophilia service.

What's your recollection, please?

A. Yes. I mean, as I say, we met on a daily basis and my recollection is that I have no recollection of requesting these results. I'd just like to enlarge upon that slightly.

> If I had requested it, it is almost certain that she would have known. I didn't plough my own furrow with regard to this. It was essential that we had this information. Within Blood Transfusion Services to this very day we still struggle about pre-test counselling. It of course -- I seem to be saying every five minutes that it was inadequate. Pre-test counselling was not the order of the day.

> We did know that samples -- I did know, personally, I did know that samples for virology were

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1 adequately. I don't recall asking for the results to 2 be there but for reasons which I can explain it was 3 essential that we had that and it was unconscionable 4 to me at the time that a family would not wish to know 5 about the health and well-being of their own children 6 and for reasons that I can explain. 7

- Q. You said pre-test counselling was not done adequately. Does it not follow from your evidence that there was no pre-test counselling at all because the samples were tested without any knowledge of parents?
- 11 A. That may be the case, although we had discussed with 12 them and there was a lot of information again in the 13 media and in the Haemophilia Society. It would be 14 absolutely -- there was an absolute need at the very 15 first moment, as there has been for every pandemic, 16 for tests to tell you that somebody has it or not. 17 The assumption which has been shown by psychologists 18 and ethicists and others, subsequently, with hindsight 19 we should have spent more time on that and done it
- better. 21 Q. Where was Dr --
- 22 A. I should have; I should have.
- 23 Q. Where was Dr Follett based?
- 24 A. Goodness. The virology reference lab. I'm sorry,

25 I don't know where it was.

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- 1 Q. Was that at Ruchill or -- it wasn't at Yorkhill, is 2 that right, or it was?
- 3 A. No, no, it was the virus reference laboratory. 4 I think it was at Ruchill.

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Q. In terms of trying to put a date on when this process was undertaken because we don't have the letter that Dr Pettigrew recalls you receiving from Dr Follett and then telling her about, we know that in December of 1984 Professor Ludlam received the information or, sorry, the information about the patients infected in Edinburgh from SNBTS product was made public in the form of newspaper articles in December of 1984 and Professor Ludlam told us about a meeting that took place on 19 December 1984 in Edinburgh.

Dr Pettigrew's recollection is that she was off on maternity leave in the first part of 1985 and returned in May and that it was in May of 1985 that the discussion took place with you about the results from Dr Follett and then the dissemination of those results. There's at least a letter from May which is from Dr Pettigrew to a doctor in Inverness with a particular result.

Are you able to assist at all with when this testing process might have been undertaken? A. I wish I could and I say that for a couple of reasons.

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Dr Gibson reports five out of ten patients testing positive for HTLV-III, which would suggest that if correct that there had been some testing by the end of November of 1984.

Do you recall whether there were -- testing was undertaken in two stages or undertaken by anyone other than Dr Follett?

A. The initial testing was done in one stage only. We did not contact, I knew Professor Tedder very well, but we did not do anything through UCL or anything like that, or Royal Infirmary or anybody else but obviously one of the issues here was is this a reliable test. It's vital, crucial, for all sorts of reasons, obviously, to know that it is and therefore it was necessary to contact the families expeditely(sic) and pass on the information that we had as well as we could and to retest and to carry out a further confirmatory test which, I believe, was western block testing at the time.

As it turned out I think that it's true to say that all of them were confirmed.

Q. The process for telling at least the current patients or their families at Yorkhill as described by Dr Pettigrew was to await their next attendance at the centre, whether it was a spontaneous attendance, as it

First of all, I don't know whether you can get more information from Professor Lowe. He may have used the same source for the testing. I don't know.

Looking at the results, there was one test in particular which was confusing because my memory, which is, I'm afraid, not precise at all, was that we heard about this at the end of 1984 and the beginning of 1985.

Amongst those tests there is you probably noticed a result from May of 1985. Now, I can't be certain about that. You can make up hypotheses but those hypotheses depend on that sample, that one sample, with no previous samples on that person, being a retrospectively tested sample, whereas it may not have been. I could only tell that if I had a letter from Dr Follett.

My memory of the letter from Dr Follett is that there were between eight and ten Yorkhill patients at the time who were infected. I'm sorry, I can't be more precise than that. You could make a theory that the results came in May of 1985. That doesn't -- that isn't my memory but my memory is far from precise.

Q. It's right to note, and you refer to it in your statement I think, that there is a meeting on 29 November 1984 attended by Dr Gibson in which

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were, because something had arisen or their next scheduled clinic attendance and Dr Pettigrew's recollection was that that wouldn't have been more than a few weeks. I think she said three to four weeks, perhaps, typically.

What can you recall about the process and your decision as to how patients should be told? A. Yes, I remember it quite well because it was a very difficult decision. To be honest with you, neither of us knew the best way to do this. I was quite sure in my own mind and -- we were both were -- about sending out a letter to people, initially with any information other than a clinic appointment was not appropriate. They were already all very worried indeed. We hoped that the majority -- and I think the majority did attend because they were mainly people requiring frequent visits to the local day care centre, that that would be the method.

And the final thing that we agreed was that we would try -- we would make sure that this was done within a six-week period maximum and preferably before. For several reasons. First of all, because we needed to pass on the information, because we needed to be sure that universal precautions were re-emphasised, and that we gave information such that

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the families were aware of, for instance, pneumocystis (unclear) VCI, because they might well require prophylaxis or immediate treatment.

So that was what we agreed and that's what we tried to put in order.

Q. The effect of doing it that way, as Dr Pettigrew explained yesterday, was that because it was often one parent only who bought the patient in, typically the boys' mother, the mother would be receiving the news of her son's infection with HTLV-III without the father or other family members being there.

Do you have any reflection or observation to make in relation to that?

A. Yes, it's far from ideal. On a balance, I think in retrospect it would have been better to bring them up to clinic visits over a period of a couple of weeks, through a letter saying something along the lines: we need to discuss your overall care and results. And such like. Or in some way that wasn't going to frighten -- I mean, getting a letter saying you have an appointment in two or three weeks to talk about something devastating is not something that is easy to live with, obviously. But with the benefit of hindsight, certainly, and probably I should have realised -- I should have realised. There was at

report, but did you have any involvement with that work?

- A. No, it was instigated by, I think, Dr Chalmers subsequently. It was a very difficult thing to do because of the sometimes patchy information.
- Q. Do you recall any discussion between you and Dr Pettigrew or, indeed, with parents, about whether to tell the boys themselves, the patients themselves, of their infection. And if so, how and when?
- A. Yes. Now, as I'd spent at least seven years in paediatrics and training as a general paediatrician before this time, the day-to-day job of a paediatrician includes talking to children. Usually with their parents, obviously, but sometimes, the older ones in particular, they were on their own. So it's part of one's job.

Now, with regard to imparting information which could be extremely frightening, especially to very young children, even as a trained, fully trained paediatrician, this is not something one should ever take on lightly. It requires specific training, specific counselling training. And one of the things that I did at Yorkhill was to bring in a social worker who had such training, and in Great Ormond Street -- and I mean training with children. And I'm very

least one case subsequently where this caused unnecessary suffering.

We always had an open door policy, such that they could come back the same day, the next day, at any time, usually Monday to Friday, but then I was there every weekend as well. So we could have dealt with it in that way, such that there was not a long delay in passing on that information. But it did cause unnecessary suffering.

Q. Professor Hann, I think the answer to this may be obvious from the evidence you have already given but it is a question I have been asked to seek clarification on.

The samples that were stored in the virology lab and presumably used for testing by Dr Follett, those would have been samples stored on a named patient basis, presumably, because the information was then coming back to you to disseminate?

- A. Yes, they were, yes, but obviously we have to rely on virologists not to disseminate information. Which they don't, in my experience.
- Q. The issue of when the patients did or may have
 seroconverted was obviously looked at for the purposes
 of the Penrose Inquiry. I'm not going to go back over
 what was said at the Penrose Inquiry or in the Penrose

pleased, by the way, that the Inquiry is looking specifically at the needs and requirements of children during that period.

The nurses that we were able to appoint were expert counsellors. We had access to expert psychology -- child psychologists and child paediatric input. So the answer is a very simple one, really, that the opportunity there for a paediatrician to speak to a child is always there. And I certainly made clear throughout my career, I hope, that that was always available. But it's a process, again. If you take leukaemia, for instance, one of the most important people is the play therapist, because they have specific training in dealing with, say, four-year old children or three-year olds or whatever it might be. How do you impart the type of information that you need to pass on to a young child who has a life-threatening disease? And the life-threatening diseases of leukaemia could mean, for instance, in acute myeloid leukaemia at the time, 70 per cent of them died.

It requires sometimes psychology and it certainly requires expert assistance. It's not something one should ever do willy-nilly, but it is there. It's not something that -- again, I really

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must emphasise, there was no question of me being the senior person there. I was responsible but we worked as a team and the team is there and is available, and to a certain extent, like the social worker, is independent of me and is available to talk to the family and the children.

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Now, final point: throughout my career, I looked after hundreds of children with leukaemia. There was always an opportunity to speak to children and we always had one mantra, which was really, first of all, whatever you tell them has to be the truth, not necessarily the whole unvarnished truth, if it's just frightening, but it really has to be given in

The second point I'd make is that, given that, and having proactively throughout my career said that, very few families actually want me, in particular -as the frightening doctor or whatever -- to talk to the children. Maybe that's a reflection on me or -but it's actually really a reflection on the fact that the others are much more experienced and skilled than me. You don't take that sort of thing with anything other than real seriousness. Children are perfectly capable, and I've seen it, of becoming seriously depressed and extremely anxious, and that can be

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had done harm unintentionally, and that probably there was very little different that we could have done about it.

That -- the Inquiry may disagree with that latter statement, and I fully accept that, but that was my transmitted information at the time.

MS RICHARDS: Sir, I note the time, and I'm going to move on to a slightly different topic after lunch.

SIR BRIAN LANGSTAFF: We will take a break in that case for an hour, bringing us back at 2 o'clock.

So time for lunch, professor.

12 A. Thank you.

13 SIR BRIAN LANGSTAFF: 2 o'clock.

(1.00 pm)

(Luncheon Adjournment)

16 (2.00 pm)

> MS RICHARDS: Professor Hann, if I might just go back briefly to one of the issues I was asking you about before lunch, which was the testing by Dr Follett.

As I understand your evidence -- please correct me if I've got this wrong -- you don't know now whether you asked Dr Follett to undertake the testing or whether he did so of his own volition?

A. I don't remember. My memory, such as it is, is that I didn't ask for that to be done.

a long-term issue.

2 Q. Did you have parents asking, on being given this 3 devastating information, how it had happened, how it 4 had happened that the treatment given by the NHS to 5 their son had infected their son with this devastating 6 disease? And if so, how did you answer as far as you 7 can recall?

A. I think I was asked this question. I think a lot of people, in Scotland anyway, in my experience -- it was 10 different to England because there was a lot more use 11 of commercial concentrates -- were wanting to blame 12 pharma basically. And the fact that we had never --13 and also the fact that the Health Service had never 14 provided enough. And, you know, we can go back to 15 self-sufficiency, if you like, but the fact was that 16 they largely blamed the commercial concentrates 17 although there was -- I can't remember the numbers but there was at least one that was probably infected with 18 19 cryo and probably one or so that was affected by SNBTS 20 product.

> The answer is that, and you have heard from others I think, that I just had to say to them, you know, we -- if we took the Hippocratic oath, we have in some ways broken it. Primum non nocere is the relevant part of the oath, first do no harm, and we

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1 Q. For the reasons we have already explored, you're not 2 able to be more precise about when that process 3 occurred?

A. No, I wish I could be. My memory, again, is that it was at the end of '84 or early '85, but there's no way of being precise unless Professor Lowe can throw further light on it.

8 Q. What would you have done with the letter from 9 Dr Follett, which we don't have -- currently at least. 10 Where would it have been placed?

A. Yes. Confidentiality was an extremely difficult issue 11 12 at this time and so it would have been in a locked 13 filing cabinet in the secretarial admin area of the 14 department.

15 Q. Do you know --

16 A. Adjacent to the laboratories, in a separate building.

17 **Q.** Do you know what happened to that letter?

18 A. I would have filed it in that way. I'm afraid --19 I took -- when I left, I thought it was inappropriate 20 to take any documentation other than very personal 21 things away with me, because they would need it in 22 future. So I left everything behind.

23 Q. You referred in your statement, and in your evidence 24 earlier, to the importance of social work and 25 psychological support for the families of the boys.

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1 We know that the social worker began at the end of 2 October 1984. What, if anything, can you recall about 3 the availability of a psychologist or some form of 4 counselling? 5

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- A. There was actually a very good professor of psychiatry, I can't remember his name, and a psychologist. I actually contacted them very early on. And also I had known a psychologist at Stirling University called Dr Markova, who was particularly 10 interested in haemophilia and the way that children 11 coped with it. I asked them if they could provide 12 support if it was required by the staff on day care, 13 and I did everything I possibly could to get expert 14 social work input, not just to this area but also to 15 the children with cancer and leukaemia and so on 16 because it was a huge deficiency in my view in the 17 department at that time.
 - Q. Does it follow, then, that in terms of any kind of counselling or psychological support, it would have required a referral to an external clinician, or perhaps internal to the hospital board, but out beyond the haemophilia centre?
 - A. Yes, there are no -- the only specific resource for the haemophilia centre was Dr Pettigrew. We had, as I say, a good but -- again, an under-resourced

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problems of families with haemophilia, including AIDS and HTLV-III antibody positivity. The Society ... turned down this application, the reason being that it was not relevant to their first priority, i.e. AIDS and HTLV-III positivity. Thus, once again, I am in the position whereby I will be financing a post directly concerned with haemophilia management, from Leukaemia Research Funds. This post will be taken up very shortly and will of course involve intensive family work."

What was the issue here? Can you recall?

- Can you give me the date again at the top?
- Q. It's 16 October 1985, I think.

Soumik --

SIR BRIAN LANGSTAFF: Yes, it is.

A. I don't have a great deal of recall of this, to be honest with you. I had been trying to throughout this period. Professor Stone I think was the professor of psychiatry. I'm not exactly sure. I think he was. And I had spoken to him numerous times about the problem not just in haemophilia but in the children and families with leukaemia and solid tumours who needed much more psychosocial support than they were receiving at the time.

Basically, by this stage, it was obvious to me,

psychology department. So we could make referrals. We did make referrals from time to time, especially in the other areas, and they were available. But it was -- really, the problem in that respect is that what you need is ongoing support. And psychologists, especially if they are very under-resourced and dealing with child protection issues, eating disorders and so on and so forth, and the death and bereavement counselling of many of my patients, they just don't have the ability to follow things up.

So they basically give diagnostic help if you think a child is becoming depressed, or if you think a parent needs help they will give -- in that era, would give diagnostic support and they always were available to speak with Anna, Sister Murphy and, later on, the social worker if a child needed specific counselling.

Q. Could we look, Soumik, at MACK0001021.

This is a letter from you to Dr Forbes in October of '85, headed "Haemophilia Counselling/AIDS and HTLV-III", and it refers to you having applied to the Haemophilia Society for a grant to pay for a clinical associate:

"The project was to investigate an interventionist approach to the long-term psychosocial

having spoken to the families in the clinic and with Dr Pettigrew reporting back to me, that there was a need for more -- as I've just described previously, more long-term involvement with the families from the psychological/social work/family therapy point of view. And although Anna was extremely good and so was Sister Murphy in their own way in providing the clinical support and social support, et cetera, I believed we needed -- and I think Dr Markova at Stirling had advised me too, having met a number of these families and looked at various problems -- that we needed long-term interventionist psychological -or at least diagnostic psychological treatment with follow-up and not just a usually one-off approach which had been available previously.

- Q. Were you able to obtain that long-term input at any point before you left to go to Great Ormond Street?
- A. I'm really sorry, I honestly can't remember. I hope so. It sounds from what I'm saying that I was intending to use private charitable funds, et cetera, for a purpose that it was never really intended, which unfortunately I had to do from time to time because of necessity. So I expect that I had done that. I think Dr Pettigrew would have known better but -- other than me, but there was certainly more psychological input

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1 from 1985 onwards. with problems but always dealt with them in the 2 2 Q. You said in your evidence to the Penrose Inquiry: context of haemophilia in general. 3 3 "We fought constant battles on behalf of our There was a lot of -- a huge amount of adverse 4 4 publicity, and the publicity was often very unhelpful. patients to prevent them being treated like lepers." 5 Could you expand upon that, please. 5 One thing that government seemed to have got a little 6 6 bit better is not putting out adverts that just A. Yes, there were a lot of good people, and 7 unfortunately a number of bad things that happened, 7 frighten people rigid. Even my own children came to 8 8 some of which I've given evidence about before, but -me one day and said they didn't want to "die of 9 9 they may seem trivial but they are incredibly ignorance", which was the main advert at the time. 10 10 important to the families. And I could list them Within the hospital itself, there was a couple 11 forever, but basically the sort of important ones were 11 of areas that were a problem. Basically, portering 12 the extreme difficulty in travel. They couldn't get 12 staff. I ended up one Easter putting children into 13 travel insurance. When they grew up, of course, they 13 radiology myself because nobody would take them. 14 couldn't get mortgages. And there was a lot of 14 There was a hospital policy promoted, which I can't 15 effort -- actually, the Haemophilia Society I think in 15 remember if it ever happened, when any child with 16 particular, and I helped them, were able to get 16 haemophilia, no matter what was wrong with them, had 17 insurances eventually. That's just one example. 17 to be reverse barrier nursed, which means they got 18 18 They couldn't get through Customs because they their food on paper trays. I even heard one person 19 had needles, et cetera, and were identified that way. 19 having it shoved under a door. 20 Within the society in general, there were major 20 But I have to say the sister on our ward was 21 problems in some schools and other schools were 21 very good in that respect and put down any 22 22 amazing. They were ostracised in schools. Some stigmatisation. 23 children had to be taken out of school or change 23 One of the very good things was our dental 24 24 department. They were marvellous, actually. I had school. It was extremely difficult. Anna and 25 Sister Murphy I know attended schools to try to deal 25 a terrible run in with a deputy chief medical officer 101 102 1 who was a public health doctor who wanted the list of 1 Covid-19, for instance, because, as you know, 2 2 our patients so that he could disseminate that to all asymptomatic transmission occurs. 3 dental surgeries so that they didn't get infected by 3 So yes, I'm sorry, another long-winded answer. 4 4 our patients. And to say that they were pariahs to There were a lot of challenges for these families. 5 5 him was an understatement. I was summoned to the Home They suffered a great deal. They were stigmatised. 6 and Health Department to -- sorry, that was a public 6 they were made into pariahs. There was good, there 7 7 health doctor. I was summoned to the Home and Health were good responses, and then eventually everybody 8 8 Department for an interview with the Deputy Chief came to their senses, but it took quite a while. 9 9 Medical Officer who said that he might have to put Q. Did you have any involvement with the transfer of 10 a black mark on my copy book, and I said he was very 10 patients to the infectious diseases unit at Ruchill 11 well -- he can do what he liked as far as I was 11 for treatment once any of the boys had become 12 concerned. It wasn't, unfortunately, in the era of 12 particularly unwell as a result of AIDS? 13 freedom of information or I might have done something 13 A. No. In fact, I don't think -- I certainly don't 14 14 remember any child becoming or developing an about it. 15 15 AIDS-related syndrome. As I've explained, I was So there were a lot of difficulties. We had to 16 16 be very careful about stigmatisation. Within the expert in the area of infection in the 17 laboratory was probably the most difficult clinical 17 immunocompromised. I had managed a lot of children 18 problem. Initially, as with, I'm afraid to say, the 18 with pneumocystis jirovecii, and I knew a lot about 19 Covid-19 outbreak, there was great overreaction. 19 prophylaxis with regard to preventing such infections. 20 I constantly reminded them that universal precautions 20 If we had got to the stage where there was an 21 in the laboratory, in home, all the rest of it, was 21 AIDS-related illness, then I would have had to refer 22 22 the patients there because there was no local -what came out -- the good that came out of the 23 23 hepatitis B tragedies. Sticking stickers all over really no local expertise. Thankfully at Great Ormond 24 samples does nothing. It does not identify who is at 24 Street the setup with infectious diseases and and

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virology and microbiology was really excellent.

risk. And it was a complete waste of time with

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Q. Prior to your departure for Great Ormond Street, had there been any advance planning for the eventuality that one or more of the boys might require admission to Ruchill?

A. No advance planning for that, because at that stage the -- we had -- I think by that stage, I can't remember exactly when, we had realised that Kaposi sarcoma and things like lichen planus and all the rest of it, wasting syndromes, were not occurring in our patients, and were less common anyway, as far as I knew, than in the other populations that have been described to that point.

So basically the planning consisted around advice about universal precautions yet again. The advice with regard to the identification of pneumocystis, which would involve persistent cough, et cetera, discussions with experts, in London usually, of when to start prophylaxis with co-trimoxazole, which had been shown to be very effective in leukaemia patients. We had a lot of leukaemia patients on that prophylaxis.

I can't remember whether or not we started those patients on it. It would have depended, to an extent, on their CD4 counts, which were done for us at the Royal Infirmary, I believe.

- hepatology is very centralised in the UK, and it's the one area where there is no major department at Great Ormond Street but there is a hepatologist within the gastro department there, so -- who I happen to know very well. So within Yorkhill we would have had to contact hepatologists in other hospitals and I would have taken advice from Gordon Lowe as to who to contact and he will have been able to give me a lot of evidence in that respect.
- Q. In relation to the period after the introduction of the SNBTS heat-treated product in December 1984, for I think about two and a half years until April of 1987, the heat-treated concentrate you would have been using for the care of patients with haemophilia A would have been NY, I understand.
- I'm sorry, I can't remember that.
- Q. In any event --
- A. The heat-treated was -- was it 68 for 24 hours?
 That's all I can remember.
 - Q. Whatever the precise name of the product, is this correct: it was believed, and we've seen your exchange with Dr Cash, to be sufficient in relation to the eradication of HTLV-III but it was understood that it did not have the same effect in relation to non-A, non-B hepatitis; was that your understanding?

- Q. In the period 1983 to 1987, was there any input from or advice sought from hepatology experts in relation to your patients?
- A. I knew the hepatologist at Great Ormond Street very
 well, Dr Peter Clayton, and I knew the infectious
 disease doctors there very well indeed, and some of
 them were my colleagues for years previously, and
 I had many discussions with them about whether there
 was anything else we should be doing, whether there
 were any tests we should be doing.

They asked me about doing liver biopsies and I -- I remember a number of conversations over whether it was worth thinking about doing any more enhanced scanning rather than liver biopsies but at that stage there was no real evidence. I don't know what happened subsequently. There was no real evidence that that helped much.

Basically, the advice was keep doing the liver function tests, seeing if they developed any other symptoms of hepatology.

- Q. What about at Yorkhill? Was there any ability toaccess hepatology?
- 23 A. That was at Yorkhill.
- 24 Q. That was while you were at Yorkhill.
- 25 A. When I went to Great Ormond Street there was --

- 1 A. It didn't have the same level of activity. I'm sorry,
 2 I can't remember what the -- whether that heat
 3 treatment had any effect at all on hepatitis C. We
 4 did know that it wasn't effective enough because there
 5 were, I believe, subsequent -- I don't know in my
 6 children's population but in other populations there
 7 had been seroconversions for hepatitis C in that era.
 - Q. Did you introduce any particular system or take any particular precautions during that period to ensure that previously untreated or minimally treated patients would not receive that concentrate and thus be exposed to non-A, non-B hepatitis unless absolutely necessary?
- A. We continued with the previous policy, as far as
 I remember, which was to use cryoprecipitate in new
 patients. As I've said, in that era it was
 a relatively small department with relatively few new
 PUPs, if you like.

There's no question in the mild or moderates.

There are very rare moderates who do bleed more, a lot more, but, as far as severes, go it would be very few new patients and they were very young, and I think

Dr Pettigrew may have told you that in any case they were usually treated with cryoprecipitate initially because home therapy in very young children -- (video

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frozen) Q. I think we lost you there for a moment, professor. The picture is occasionally breaking up. Would you be able to repeat that last answer you gave. You said home therapy in very young children ...? A. Oh sorry. In very young children, in any case, cryoprecipitate was usually the standard of care in that era unless there was a severe episode like a bad nasty head injury or whatever. But as you probably know, in the first couple of years of life such episodes are uncommon anyway. Many of them don't even present for the first year or so of life. But if it was required to maintain definite haemostasis and for that to be continued over a period of time, then cryoprecipitate would not have been a good solution for several reasons, not least because even during this period we were still having some difficulties in

Q. Can you recall what efforts were made to address the possibility of infection with HIV for haemophilia B patients during the period up until I think October 1985 when a heat treated Factor IX product became available from SNBTS?

of fears over cross-infection.

getting factor assays done in the laboratory because

A. Gosh, this is stretching my memory. We had very few

the first half of the 1980s at Great Ormond Street?

A. Well, I was very -- yes, during that time. Basically, when I was a lecturer, I was there, you have the dates and I've forgotten them again, it was prior to 1979, so it was about 1977, the patients that I saw were almost all on concentrate and a very high proportion of that was commercial. I think a lot of it was Armour. That's just a memory but certainly there were -- there were no patients that I met during that period of time that were on home therapy with cryoprecipitate.

Again, there was no written protocol at that time when I was a lecturer. Sorry, are you talking about 1988 now?

- Q. I am but I was asking what you knew of what happened before and by all means fill in that gap by reference to your own direct knowledge from working there too.
- A. I was there for two years so I can remember a fair bit of it.

Cryoprecipitate, there was no written protocol. There was an understanding that patients with mild haemophilia and von Willebrand's disease would be treated with cryoprecipitate. There was a very small amount of use of DDAVP. We used quite a lot of antifibrinolytic agents, tranexamic acid, EDHCA,

patients by chance with Christmas disease, haemophilia B. I honestly cannot remember. It may be one or two. In those patients the only alternative is fresh frozen plasma which is even more difficult than using cryoprecipitate because the amount of Factor IX that you get in that is minuscule compared to the concentrate.

So basically again the idea -- and it also doesn't respond to either cryo or DDAVP or whatever -- so basically if there had been, and I don't remember any episodes to be honest, where it also tends to be -- there is a tendency for it to be a milder clinical manifestation as well -- not always but it quite often is. My memory, such as it is, is that we rarely needed to treat patients with Christmas disease and that when we did, we would have tried to avoid the use of concentrate during this period of time where possible and even consider the use of fresh frozen plasma, although it was a very poor treatment.

Q. I want to move next to some questions about Great Ormond Street. When you arrived, the concentrates that would have been available would have been I think entirely heat-treated.

What was your understanding or what did you learn about what the approach to treatment had been in

et cetera. For patients with Christmas disease, they were treated, as far as I remember, exclusively with fresh frozen plasma and the patients with severe haemophilia were, by that stage, the vast majority of them, even though they had only been there for a year or so, were transitioning to home therapy and eventually the idea, as in Glasgow, was that they would transition to adult care once they were self-treating and established in secondary school at about the age of 12 or 13.

- Q. When you returned to Great Ormond Street as a consultant following your four or so years at Yorkhill, what, if anything, did you understand or learn to be the position about the circumstances in which Great Ormond Street patients had been infected?
 - A. Well, it had been managed by Professor Hardisty and one of the lecturers, Dr Ball, mainly in collaboration with the infectious disease department and one of the lecturers there who was given that as a specific role.

There were quite a large cohort of patients, I don't know the number I'm afraid, who were infected with HIV, some of whom -- a few of whom had developed problems, one of whom I remember had developed autoimmune thrombocytopenia with a low platelet count, but there weren't a great number of patients, very few

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in fact, who were developing overt signs of an AIDS-related type syndrome.

Where that happened the studies had already been undertaken and were in progress with AZT, which was led by the infectious disease department and the professor of immunology, Professor Levinsky.

- Q. As and when patients of Great Ormond Street did develop AIDS and opportunistic infections, how was their care and treatment managed? Were there joint clinics or was it a question of a referral to the infectious diseases unit?
- A. I didn't really run joint clinics because yet again I was pretty much on my own. What we did was they would be seen in an infectious disease clinic and then I would see them in a clinic, either the haemophilia clinic or in the haemophilia centre, which was very limited initially but at least it existed slightly.

The management of these patients was essentially by the infectious disease and immunology team with the hepatology input but also we relied a very great deal on to our clinical nurse specialists with specific children's counselling training in haemophilia and infectious diseases and the infectious disease lecturer who -- all of whom, if you like, co-ordinated as a team.

to every patient who was about to be tested whether or not they turned out to be positive.

- Q. What, if any, role did you have in the treatment of patients with bleeding disorders who had been infected with hepatitis C as and when treatments became available in the '90s, interferon, interferon and ribavirin and so on?
- A. As far as I remember, as I've said, I think there was only one patient between '88 and 2006 who required interferon therapy. It was extremely poorly tolerated. It was not -- I didn't -- I always believed that specialisation and expertise was important and I certainly didn't dabble in infectious disease management or HIV management or hepatitis C management other than looking after their haemophilia and ensuring that the co-ordination between the multidisciplinary approach worked, which was also particularly the role of the nurses involved who worked together.
- Q. At what age typically did the bleeding disorder patients at Great Ormond Street cease to be the responsibility of Great Ormond Street and be referred to an adult service?
- A. It depended on what was wrong with them. If they were not infected, they were usually transferred and again

They also had a lot of input from psychology and social work and it was a very well resourced area.

- Q. When the test for hepatitis C became available, what can you recall about the process for testing patients for hepatitis C at Great Ormond Street, the bleeding disorder patients?
- A. Thankfully it was a great deal better. Pre-test counselling was intensive and expert from the -- I think there was eventually even a hepatitis C clinical nurse specialist but initially it may well have been from -- yes in fact there was, from the infectious disease team and, to a certain extent, ourselves because this was across the board. There were -- being a tertiary quaternary referral hospital there was a lot of patients who received a lot of blood products for cardiac disease, for immuno-deficiencies, for all sorts of other things that were not related to haemophilia, and of course all the leukaemia patients were at very significant risk.

So it was a very big undertaking. It again required retesting and gold standard, as it was called, testing but it was -- by this stage, I suppose we'd learnt our lesson and pre-test counselling and post test counselling and ongoing support was provided

adolescent management in the UK has never been brilliant. It's a great deal better now. We've worked very hard and we then my successor, Dr Liesner, my colleague and successor, managed to set up a transitional care consultant between ourselves and the Royal Free and some of the patients south of the river went to St Thomas'. So the answer to your question is that we would expect them to be on self-therapy and established in secondary school. That establishment was managed from, as an outreach, from the department.

Now, if they were infected with hepatitis C or HIV or both and had no problems whatsoever and were not requiring any treatment, and I don't know if there were any of those, they would also transfer to those centres with the transitional care put in place.

Those who -- those patients who sadly were running into major problems actually stayed between ourselves and UCL which there was an -- a lot of us had joint contracts with UCL. The infectious disease people, the immunologists, some of them did, and they managed the patient between ourselves and UCL.

We weren't allowed to admit patients over the age of 15 to Great Ormond Street except in exceptional circumstances, so it was essential that there was some

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sort of joint management between those two centres, because they had a great deal -- you know, that's where Dr Tedder worked and, you know, there was a great deal of expertise there, in HIV and hepatitis C, and they basically managed the patients. As I say, it would have been foolish for me to

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interfere with that but we made sure their haemophilia was also managed and they weren't stigmatised in any

Q. Professor Hann, I have some more general questions for you now, not specific to Yorkhill or Great Ormond Street but to be answered as you're best able to.

Did you consider that the management of children with bleeding disorders gave rise to different ethical considerations than the treatment of adults and, if so, in what respects?

A. Yes, absolutely. I suppose I would say that, wouldn't I? I'm a paediatrician. Children are not small adults. They've different disorders, diseases, they have different responses to things. They have much, much different educational needs. They have different psychological needs. They need expertise.

This was a question that arose and I won't ramble on, but this is a question that arose repeatedly within this particular area which was

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of that. So I do think that children should be managed within centres where there is a lot of hands-on paediatric experience, not just the great almighty doctor but the whole team and what we used to call the seamless roving in Great Ormond Street, in other words, everything around it that's required for dealing with children with long-term disease.

That finally by the way doesn't mean that we had to do everything. We have, for instance, a hub and spoke approach towards leukaemia treatment, et cetera, and we empower centres like Watford, or Welling, or the Whittington or whatever to manage, co-manage those patients with us. That's more difficult to arrange with haemophilia because it's far less common -- it's less common and much more chronic. But anyway sorry that's a long answer.

- Q. Are there, in your view, any different ethical considerations that might arise as between the position of children and adults when it comes to treatment that might be regarded as experimental or particularly high risk?
- A. Yes, there are, and I've already explained to you that -- the involvement of people like play therapists and all the rest of it, explaining -- it's a question of explanation, appropriate explanation, an

almost unique in medicine in that many children were looked at after within adult centres with adult physicians looking after them, maybe having some advice from paediatrics.

In Glasgow that was not the case and latterly although only quite a lot latterly that was -- it was the case in London that a lot of children were looked after in major adult centres by adult-treating physicians.

It would virtually be the story of my life if you like if I were to say what is different about paediatrics. Does it matter that you are paediatrically trained? Yes, it does. Does it matter that you are in a centre that has multiple paediatric specialists and sub-specialists? I believe so, which is one of the reasons why I moved. Does it matter that you have input from people who are trained in both counselling and communication and all the rest? That's not just psychologists, the nurses in particular and the social workers with family therapy counselling training are a vital element here too.

So they also obviously do have some different diseases. They have different ways they present. Children with pneumocystis do present somewhat differently. For instance, I had a lot of experience

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explanation that doesn't just frighten but informs appropriately and honestly and transparently.

The ethical considerations are -- I've read the ethical report, which is very good. The problem with ethics when you're on the ground is it's often very difficult to decide what is ethical and what is not. The legal aspect of it really isn't the major issue, although it was clarified with Gillett principles and so on. I was very unhappy prior to that that children were assumed not to be able to give consent to anything.

There are episodes, and I give you the example of -- I know it's not the same, but children with bone tumours, for instance, where amputation is an issue, and the child may be 10 or 11, or whatever it may be, these are difficult -- and may refuse therapy. Do you override that or do you regard them as being competent or do you -- what do you do?

We set up a forum. We had obviously an ethics -- not department, but we had a paediatrician who had trained with Sir Ian Kennedy, who you may know, and who gave us advice. And we also had an ethical forum, for instance, for children who had multiple congenital abnormalities, and of course

that's been in the press in the near past (unclear).

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These are very, very difficult issues sometimes, where parents expect treatment that doctors believe to be futile, where children refuse treatment and the parents want it, where -- you know, all of these things come up, and are not theories, they are actual practical problems.

I was very lucky at Great Ormond Street, we had usually very good advice, and we're surrounded by people who had the best needs of the children in view.

- Q. In your witness statement you made an observation that blood transfusion centres in England seemed to function like individual fiefdoms. What was the basis for that view and are you able to elaborate upon what you meant?
- A. Yes. There is no doubt that's how it worked, in my view anyway, when I was training, and that would be in the mid-1970s.

It was set up, like a lot of things in the NHS, piecemeal post-war, based on people who had been in the army and places that had been set up to provide blood and such like, not necessarily in the right place and not necessarily with all the best people involved, not necessarily, or, you know, all the appropriate expertise.

I don't know when this changed in the UK. It

"Training of all clinical staff with regard to blood and blood products and [transfusion transmitted infection] is still not satisfactory ..."

And you say:

"For some reason this area is still not taken seriously enough and mistakes such as wrong and inappropriate transfusions continue."

Could you elaborate upon that, please.

A. Yes. It's obviously improved and we're, within transfusion, doing everything we possibly can to ensure that -- for instance, within Ireland we now insist that all trainee staff when they come in have carried out the E-learning facility and been accredited on it before they come.

We still have to put a great deal of effort in by our -- one of the great things that came out of this tragedy is that haemovigilance and governance of blood products improved out of all proportion with the appointment in every hospital of haemovigilance staff, who report directly to the hospital transfusion committee and the executive team.

So it's unfortunately the case that within medical education in the UK I can tell you, whatever those educators say, when they come to the hospitals their attention to detail and their knowledge is often

certainly had changed by the time I got back to
Great Ormond Street. You know NHS BT is a world-class
institution, with fantastic scientific backup, which
we rely on a great deal in Ireland, by the way, even
after Brexit, and they work as a whole team across the
country now.

For instance, you know, they have centralised things like frozen storage of rare blood groups in Liverpool, and they have groups of scientists and technicians and laboratory scientists who advise them on things like blood products, radiation and particularly, the foremost fear still, transfusion-transmitted infection, which is a day-to-day issue still within blood transfusion.

So in those days it was down to the local director. It was run by a doctor who was sort of all-seeing and all-pervasive and who made the decisions and who seemed to run the budget and everything else. So it was pretty much a fiefdom.

There were major problems in the late 1970s because of activity and it came to be carried out in the courts subsequently, and I think that was one of the drivers to put it right.

Q. You identified also in your witness statement your concern that:

initially not sufficient. And I am not talking about haematologists here, because that's rammed into them, but in general there are still some doctors out there who regard a bag of blood as something that sticks your haemoglobin up by a gram or so, if you're an adult, and that's about it, despite the fact that we've spent forever producing leaflets for patients and getting consent organised, et cetera, for every transfusion.

Q. In your evidence to the Penrose Inquiry you described this:

"There was inappropriate use of blood products on a regular basis, particularly by surgeons, who often attributed near magical properties to products such as fresh frozen plasma and cryoprecipitate and whole blood."

Were you talking about a particular time period then, or is that something that you've been aware of throughout your career?

A. There is still -- it's improved a great deal. There was a lot of belief. It was a bit like phlogiston or -- anyway, it was a pseudoscience-type belief in things like whole blood: it gave you everything you needed bar the kitchen sink, it stopped you bleeding, it plugged up holes, it stopped surgical bleeding, it

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important actually.

might even have treated infection or whatever.

As I say, the education in those days was wholly inadequate. It has improved a lot but there are still -- there is still inappropriate use from time to time, which is monitored through the Blood Transfusion Service attending hospital transfusion committees through IRs, as they are called, incident reports, which every hospital in Ireland anyway has to produce.

We look at things like why are people still using cryoprecipitate when it is no longer the safest drug for surgical bleeding following cardiac surgery, for instance. Why are they still doing it? And the evidence that exists -- and unfortunately, one thing I will finally say is that I grew up throughout the 1970s and onwards with the management of children with leukaemia. It's been a great success story and it's based on proper data, proper clinical trials, proper outcome, proper governance and overseeing of those results. Blood transfusion did not work like that. has not worked like that, and the number of clinical trials, randomised trials, carried out in this area is still far too small. This has been recognised by the Cochrane reviews, et cetera. I recently looked at two things: like screening for CMV negativity in blood

treating inhibitors. It's absolutely massive. And it's a massive cost as well, of the order of something like half a million per year's treatment.

So you could say well neither of these things were established. Well, in actual fact, the evidence from the 1970s in Sweden were fairly compelling, and we were able to confirm -- or Dr Liesner was able to confirm that, even in patients who had very bad target joints.

The evidence with regard to inhibitors came from Germany, which has always done much better than us in this area, where they were treating -- I think they were called Bonn Protocols and so on. They were treating patients with very high dose so-called immuno-tolerising treatments well before we were able to fund that or get enough treatment in the UK.

Things have changed radically. But actually being at a full level of self-sufficiency, we were a very long way off that. Scotland was much closer and probably could have taken on prophylaxis for the majority of children eventually, but obviously the finding and verification of recombinant products solved the problem. If you could afford it.

Q. You also refer in your statement to two concerns: one, the lack of a national expert advisory body or group products, since the 1960s and '70s there have been ten trials only; irradiation of blood products for patients who are having bone marrow transplant, zero trials. It's still a bit of a Cinderella area of medicine.

- Q. You have referred in the course of your evidence today on a couple of occasions to the issue of self-sufficiency and wondered whether we would go back to it, and this is my invitation to you, Professor Hann, to go back to the issue. You deal with it in your statement and talk about the fight for self-sufficiency. What would you wish to say on that issue?
 - A. I won't keep you too long on that because this may well be irrelevant.

Self-sufficiency means that you can treat every patient optimally. That would mean being able to treat everybody with prophylaxis, which is a huge extra drain on resources, and also as big a problem, being able to treat the inhibitor patients with massive doses of Factor VIII. There was no way that the UK was in a position -- or appeared to have been unable to be in a position to get to that level.

I can't remember the doses that you would require for prophylaxis per year but it's greatly increased for

of some kind during the 70s or 80s and, secondly, the lack of national guidance.

Again, can I invite you to elaborate upon how, in practical terms, that could have assisted?

A. I'm very grateful to you for asking me that, genuinely, because it was -- you know, if nothing else good should come out of a tragedy like this -- and I've already mentioned some of the good things: the governance of blood products, the traceability, which is a legal requirement in Ireland now. But there are some things that have not happened. I was going to say this in a concluding statement but in a few minutes I would like to address it now because I think it's very, very -- I think it's fundamentally

This was an era where we were coming -I believe coming out of a very patronising era in
medicine. There were still some doctors around who
believed that they were some sort of god or whatever,
but not within my haemophilia community. What we
needed was not a bunch of doctors meeting together
with a bit of advice from somebody who had some
interest in some area related to what we were dealing
with. What we needed -- and this was not solved with
hepatitis C entirely and certainly it was not even

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answered when we came to prion disease, and I said that in my statement. What we needed was -- and I'm afraid to say -- it has to be either governmental or at least it has to be in a position to give you something in authority which you know you can follow, with knowing that it is correct and it is coming from the right sources, et cetera.

Basically, I would describe the situation that we were in was in some sort of maelstrom, in a whirlwind of different advice based on very little science, coming at you from all different corners, and consensus very difficult to achieve in certain areas, also including non-A, non-B hepatitis. It was very difficult but it would have been a great deal better if there had been central guidance, if we had had some sort of SAGE committee or whatever at that time. I'm sure that's much better now. I sincerely hope it.

But it's taken this pandemic for this to improve and I'd just like to say two other things really because I think it really is of fundamental importance and I will be quiet after that.

One of the major problems we had at that time was with public health input. Some people will hate me saying this but the fact is that it was either not much help or it was actually unhelpful. What we

are going to start taking viral infections and transmitted infections seriously immediately or very soon has been a very, very disappointing process. Most virologists now accept that we are going to invariably suffer such plagues and we need to be better prepared. This is an existential threat to humanity, possibly even as serious as that of climate change.

We've already in the Blood Transfusion Services had to deal in the Americas with Zika virus causing major neonatal problems, with West Nile virus, for which we have to test in Ireland at the moment, and many other problems such as, for instance, hepatitis E, which is very common in Ireland and the UK at the moment and is associated with a degree of uncertainty.

Worse than that, there are other viruses coming through which get very little publicity. For instance, Nipah virus has already killed at least 300 people in Malaysia. It comes from bats, through pigs. I don't think anybody else ever again is going to really look at porcine products for human use. Hendra virus has killed a number of people in Australia and comes from fruit bats, which really takes me on to the final point that I wanted to say as

needed from public health was experts who had real experience of dealing with public health issues and not just preventative health ideas, which obviously is a very important part of public health, but actually real training and real experience of dealing with epidemics, pandemics, endemics, whatever you like to call them, real experience. My experience subsequently is that this has improved but it's still not adequate.

You know, you only have to look at Ireland, where I work now. Public health doctors are not even recognised as consultants and are not paid as such. It's a disgrace. It's been put forward and brought forward so many times. You know, if you ignore history, you are undoubtedly doomed to repeat it. I think that training in those areas still has to be addressed, it has to be addressed with proper hands-on training. There are people, Dr Markova and various others who are very experienced, one of the leaders in the WHO is a Dr Ryan, an Irish man.

So the expertise does exist but it does not exist across the board to a level which would have helped us at the time or subsequently.

The other thing that I really wanted to say was that to come out of this and to believe that people

a conclusion which is that, you know, this is an area that should be foremost in research funding in Government priorities and in all aspects of medical training.

Also, that research needs to be between the veterinary and the medical fraternities. There's very little — there is some but there's very little collaboration at this moment in time. That must be improved. Experts in zoonosis need to be part of the advice that you get during these epidemics along with behavioural scientists, psychologists and all the other things that we didn't have at the time.

We still haven't learned all of that. And until we do, we are going to have further problems, which are going to be a major threat. And just one final thing, if people don't think research is impossible, just think for a moment: how is it -- if the bats are the carriers and live with these viruses within them and have methods of control of these viruses within them, that should be a very, very important research protocol within any country at this moment in time.

So, I'm sorry, I just wanted to take the opportunity to say that there were -- this was a terrible tragedy, nothing worse in my experience of

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1 over a third of a century of treating children, but we 2 had hoped that good would come out of it. A lot of 3 good did come out but there are still major things that we have not learnt and I hope that this Inquiry 4 5 will bring to the forefront. 6 Thank you very much for allowing me to say 7 that. 8 MS RICHARDS: Professor Hann, sir, I have got some 9 specific questions that have been suggested to me by 10 recognised legal representatives of Core Participants but we need, obviously, to give the opportunity to 11 12 them to suggest any further questions. I don't 13 anticipate there will be a huge number more but, 14 bearing in mind the need to give that opportunity and 15 the fact that the picture is breaking up somewhat from 16 time to time, I wonder whether we could take the break 17 now and then when we come back I can finish the

SIR BRIAN LANGSTAFF: Yes, indeed. 3.30. Take a break until 3.30, professor.

questioning of Professor Hann by reference to the

further suggestions I might receive from Core

23 A. Thank you.

Participants.

24 (3.00 pm)

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(A short break)

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arrived? Was it the Health Board, Dr Willoughby, the Scottish Home and Health Department, or a combination?

A. Difficult to say. All I know is that when I came there, there was a lot of -- there were a lot of people who were trying to help me improve things, including people in the Health Board and all of the areas that you mentioned.

I think Dr Willoughby was possibly even worse at politics than me. He -- I can only pass on things that other people -- hearsay from other people really, to be honest with you, in that they said that he didn't fight hard for things and such like. I really, honestly, don't know, but it's important to understand two things.

First of all, this was -- paediatric haematology and looking after children with leukaemia and blood disorders, bleeding disorders, was a relatively new thing. There were a lot of very poorly resourced departments, including Great Ormond Street at the time. There was half a haemophilia nurse when I went there and a store cupboard converted into a bleeding room and an area you could barely get in through the door where you treated patients.

When we started doing external audits, my first visit was to Belfast, to the unit there, some years

(3.32 pm)

MS RICHARDS: Professor Hann, a number of further
 questions suggested by core participants that I'm
 going to ask you about.

In terms of the facilities at Yorkhill on your arrival, what were the available laboratory facilities and where was the testing on factor levels and ALT levels undertaken?

A. Basically there was a -- right next to my office there was a main laboratory, and then next to it there was a blood transfusion area. That was it.

The clotting initially -- coagulation testing -- was done in the main laboratory, and then I was able to send, with support, I think mainly from charity again, a lady to Canada, because Dr Gibson had worked in McMaster and was able to get additional training for that person, and she came back and ran the clotting lab thereafter.

- Q. So in terms of testing for ALT levels, was that undertaken which you have described in Yorkhill or was that sent off elsewhere?
- A. No, that was just down the corridor, in biochemistry.Dr Logan I think it was.
- Q. Who would you say was responsible for the lack offacilities and staff which you observed when you

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later, in around about 1990, probably, to Dr Elizabeth Mayne, and there was effectively no haemophilia centre there. It was a matter of using beds on the ward or maybe areas in clinic, et cetera. It was a common problem. As I say, although things were designated as centres, they weren't resourced as centres. There were exceptions to that, and the best exception by a long -- actually Royal Manchester Children's and Alder Hey did not have any designated centre area either, when I've been there very recently. The best example of a well-resourced centre, and one of the very few at the time, was the Royal Free, the Katharine Dormandy Centre, which I think was started in a caravan, but with the rebuilding of the hospital in Hampstead, as some of you may know, it was all-singing and all-dancing.

Sorry, as far as staff were concerned, yes, it was -- I think that there was -- I've already said there was a lack of sub-specialisation in paediatrics in Scotland at the time. It was not given a high priority really by anyone until Dr Willoughby left and I came, and his leaving was part of the drive towards this. There were a whole series of appointments in paediatrics and areas of general paediatrics and neonatology and so on, yet I couldn't get a paediatric

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oncologist on board. I did manage to get Dr Gibson there, which was a godsend, and she was extremely good. But getting other things, like facilities, I had to raise the money myself.

Q. You described communication issues as the biggest

- Q. You described communication issues as the biggest problem on your arrival. How did those problems manifest themselves?
- A. Let me put it this way, all the things I was used to -- of all the things I was used to, very little existed. I was used to social workers with family therapy training, I was used to clinical nurse specialists who often had counselling training, I was used to the availability of psychologists to a somewhat limited degree. And I had, as a trainee, significantly more time available and personnel available to carry out communications. But in addition to that, I noticed that a number of families were not part of The Haemophilia Society and didn't seem to know it existed and I had to push that very hard.

I was very used, at the Royal Free, to a routine clinic. In fact, there were clinics every day there virtually. So that was something that needed to be sorted out immediately. Then at Great Ormond Street in particular, through the palliative

et cetera, and you know how dare they step on our toes as nurses and doctors and so on. We are there and we understand how to talk to people and how to communicate and all that, not realising that there is an absolute need, especially in this sort of area and especially when there's a lot of grievance and grief and so on, for somebody independent with proper skills to be available to them, so I'm sorry I'm being very verbose but it was something that I had to fight for very hard.

Fighting for resources is one thing and I wasn't all that good at that. Fighting for a new ethos is even more difficult, but with the appointment of a very good social worker for the leukaemia solid tumours, et cetera, and the involvement of Ms Leitch, it improved dramatically. I think her input as what I would call a patient advocate is, especially in times of crisis, is so important.

- **Q.** How did parents react to the changes you implemented in terms of the switch to SNBTS concentrate and the cessation of prophylaxis?
- A. There was -- I mean, I have to be honest with you in that this was dealt with almost entirely by Anna Pettigrew and Sister Murphy, and they fed back to me and I did talk to them in the clinic about it and so

care and home care teams, I was used to the communication -- the excellent communication skills of proper paediatric trained nurses and nurse counsellors through parents' groups, which were either *ad hoc* on the ward or whatever, but very often, for instance, bereavement counselling sessions and information sessions. Like, for instance, when there was a new leukaemia trial, et cetera, there would be open sessions, town hall-type sessions.

Also, on top of all that, there was an absolute need, which had not been recognised at the time, for independent patient advocacy, if you like. Something that came later on at Great Ormond Street that was actually just inside the main door of the hospital, probably still there, was the patient advocacy team which I fought for with many other people and which reported directly to the Chief Executive and to which people could drop in and was entirely confidential and also the extremely important role of social workers.

I will just add one thing, and this is not intended to be pejorative but the attitude towards social work at the time needed a bit of a revolution in Yorkhill and the attitude basically was these people were there to sort out a bit of extra money to buy fridges and freezers, a bit of extra clothing,

on.

There was a little bit of a reaction because you know the marketing of the product is obviously as you would expect better with the company, nice packs and bits and pieces and swabs and butterflies and all the rest it. I have to say their product information, or PILs as we call them nowadays, leaflets within the packs which actually dealt from the mid to late 1970s quite well with the risk of hepatitis being there.

All of those things were preferable. It was of higher purity and they knew that.

But basically I just said to them that I can't prove it to you but, you know, it's almost common sense that the approach of, in most respects anyway, of the SNBTS was -- and the blood Transfusion Service throughout the UK was preferable.

- Q. One of the aspects of your policy changes was to try and keep concentrate use to a minimum. How did you control concentrate use amongst those who were on home treatment?
- A. I didn't actually really control it that much. The only thing that I did or two things really was that I reduced -- it was very important to me and to the country at the time that we preserved the supply as much as possible so that we didn't have to go back to

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the use of commercial concentrate. So I did -- I had two reasons to stop prophylaxis. First of all, it wasn't working in the way it was being given to those patients who were on it and, secondly, with patients who were on home therapy and so on it was a routine anyway to be sure that it was being used in not a profligate fashion. That very, very rarely happened anyway.

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So we didn't say to people you can't do this anymore. What we said to them was, "Prophylaxis isn't working. We do need to maintain the supply. I would advise you that we should cease prophylaxis until we know more about how it should work", or whatever, and I don't recall any major push back to that, although one or two people were a bit fed up with having to spend longer, it's just a few minutes longer, drawing up the product which was of lower purity from the SNBTS.

- Q. Were you aware from your work in the field of leukaemia of the work in the very early 1980s of the Gallo group on HTLV-I and II?
- A. I had actually met Bob Gallo at various meetings and
 so on. I knew his main interest was, to my knowledge,
 in bone marrow transplant and so on. But I did I may have been aware of it. It wasn't something

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- protocols of that nature. It would have been along the lines of we need to do this because the patient might be considered for pneumocystis prophylaxis, for instance.
- Q. Was any research undertaken on your bleeding disorder patients during your time at Yorkhill?
- A. Yes. I've already mentioned Dr Markova in Stirling who had had a long -- prior to my being there, I think, had had an interest, and I encouraged that interest because we needed more psychological, psychotherapeutic sometimes, input. Her interest was mainly in coping procedures and -- in children with haemophilia, things like, you know, what could they do with regard to sports, were they scared to death of knives and and so on and so forth, how do they manage scratches and how -- how are they adapting to their life with haemophilia. And that was done, as far as I'm aware, with the full consent of those taking part. I mean, she was a psychologist (unclear) and she was a very good psychologist, she published quite a bit, which some haemophilia families read of course. And I'm sure she explained it in a consenting way.
- Q. Was there any other research in which the patients were involved, to your knowledge?
- A. I can't think of anything else, no.

1 that -- HTLV which did you say?

2 Q. I and II.

A. HTLV is still a problem to this day as
 a T-lymphotropic virus. I didn't actually know that
 he was particularly interested in it but I was
 certainly aware of HTLV as an issue in leukaemia
 patients and it's really a problem arising, my memory
 is, out of the West Indian population and so on.

- 9 Q. In the period when you were at Yorkhill before you
 10 learnt of the HIV infections in your patients, were
 11 your patients tested for immune function deficiency?
 12 Were CD4:CD8 counts undertaken prior to their
 13 diagnosis?
- A. I don't think so. I was very much aware of one issue which I regarded as a somewhat ethical issue, which is that it was -- I don't think it was suggested in
 Scotland, but it was certainly suggested at some of the meetings that I was that one could surrogately test for HIV by doing CD4 counts. I did not approve of that approach.

If there was anyone who had, and there were not to my knowledge, if there were anyone who had shown any signs of AIDS-related illnesses of any description we would certainly have done it then. I don't think we did, no. I certainly didn't have any research

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- 1 Q. Just going back to the issue of the letter from
 2 Dr Follett and the process of testing for HTLV-III, do
 3 you have any recollection as to whether that letter
 4 contained only the results of patients who were still
 5 Yorkhill patients or did it also include patients who
 6 by that time had moved to the Royal Infirmary in
 7 Glasgow?
- 8 A. This really was why I pointed to Dr Lowe may be able 9 to help more. As far as I recall, and I think I read 10 some of Dr Pettigrew's statement, that it was the 11 same, that there were about eight or ten names on that 12 list. I'm as sure as I can be but I'm not absolutely 13 certain that there are no transfer patients' names on 14 that list. The fact that they were done at the same 15 time may give you a clue, with Dr Lowe, as to when 16 those tests were actually performed. 17
 - Q. Did Dr Gibson have any involvement, as far as you can recall, with the process of testing?
- 19 A. I very much doubt it. I don't think so.
- Q. You referred I think in one of your answers earlier to
 the possibility of a patient having been infected with
 HIV from cryoprecipitate. Are you talking there about
 a Yorkhill patient or elsewhere?
- A. I'm sorry, I don't have the list in front of me.
 Within my statement I just reiterated what Dr Chalmers

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- 1 did. It's in black and white. I never did what
 2 I know Sir Brian was describing and what we still call
 3 a look-back. I never did that. But my memory is that
 4 there was a patient who developed HIV from
 5 cryoprecipitate.
- Q. But you have taken that simply from the work recordedin the Penrose report; is that right?
- 8 A. Correct. From that table, you know, the one you --
 - Q. Yes.

- A. It's actually -- and there's a sort of diatribe that
 goes with it as well, yes, which describes where it
 was possible to try to elucidate whether it was SNBTS
 or commercial or cryoprecipitate as stated, and I put
 it in my statement.
- Q. What, if any, psychological or other support was
 provided specifically to the parents of the children
 infected at Yorkhill?
 - A. Well, we've already stated the difficulty that there was with mothers being involved initially, but we always encouraged them, if possible, to involve the other parent or partner. When it came to the parents' support groups, everyone was invited. There was never any question of identifying people who were positive or not positive. It was generic, initially, and then I believe that the social worker and the parents

there sometimes are. So -- and we still worry about
that side of things in blood transfusion. How long do
you keep Health Life questionnaires? 10, 20, 30?
Simple laws don't really help you in those
circumstances always.

- Q. In the period from the end of 1984 through to April 1987, so the period when the SNBTS product protected in relation to HIV but not in relation to non-A, non-B hepatitis, were you aware of there being, in Scotland, any supply of 8Y, which was the BPL the Elstree-produced product?
- A. I know it quite well because I obviously used it and did so afterwards. I don't recall ever using 8Y in Scotland. It was a very good product in many respects, had a very low inhibitor risk apparently, but I don't recall using it in Scotland.
 - Q. Final question, Professor Hann. You referred in your evidence, I think this morning, to UKHCDO as a doctors-only organisation. Who else do you think could or should have been involved to broaden it out?
 - A. Yes, well, I was a bit of a lone voice with this.

 I probably might well still be. I believe very strongly that most of our haemophilia care should be nurse-led. It's always, from the mid-1970s, been a home care type of approach, a day care type of

themselves held parents' groups themselves.

We certainly, and Dr Pettigrew -- I spoke to one group and I know Dr Pettigrew at least spoke to one group, if not more. It's possible that the invitations didn't go out to everyone, and that's regrettable, but I think that we made every effort, and the social worker and others certainly made every effort to get everyone there that could be, available.

- Q. Dr Pettigrew's evidence in relation to patient records was that records were retained at Yorkhill when a patient was transferred to the Royal Infirmary in Glasgow. Is that your recollection and if so what was the purpose of that?
- A. I'm not sure that there was a specific purpose.

 Basically it was the case, yes. We occasionally loaned out case notes, but sometimes, for all sorts of reasons, you get contacted at a later stage about somebody asking for information about a patient, obviously in a confidential manner. So we -- my attitude was I sort of ignored the 25-year rules and and so on and I always said, both in Yorkhill and at Great Ormond Street, that we should retain the records of these patients forever. That wasn't always followed, of course, but -- and that causes problems when there are things many years down the line, as

approach, co-ordination of such patients, doctors are very bad at that on the whole. The nurses have nurse counselling expertise or should have and I really thought that at the very least it should be broadened out to include them.

I don't think there's any specific reason why that view only have a doctors' only -- well, originally it was only reference doctors. It then became broadened out to more doctors. Certainly the annual general meeting could have been for nurses and I would just reiterate that there was one thing which was remarkable, which was that the World Federation of haematology was one of the very first meetings ever where everyone, including the patients, was invited, and they took a very active part in it and, by the way, were supported by the commercial pharma companies to do so.

So, yes, you could say everybody and maybe there should even be a national UKHCDO which includes patients but, as far as the planning goes, it would certainly at least be important to include nursing and you could also consider including physiotherapists and social workers up to a point. I mean, sometimes when you're dealing with very technical issues you have, as the World Federation does, you have break-that type of

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1 approaches which are intended to be more of interest something to improve, and that's what happened with 2 2 as opposed to keeping people out. So, yes, sorry, HIV. It happen again with Zika virus such that the 3 3 long answer again. Americas, USA particularly, introduced pathogen 4 MS RICHARDS: Professor Hann those are my questions and 4 reduction techniques for cellular products which are 5 I understand that your counsel, Mr Bowie, doesn't have 5 much more difficult to deal with than plasma products. 6 6 any questions for you. So Ireland, for instance, and the UK are 7 Sir, over to you. 7 looking again and we hope to introduce very soon 8 8 Questioned by SIR BRIAN LANGSTAFF pathogen reduction techniques for platelets and then 9 SIR BRIAN LANGSTAFF: Yes. When you were answering 9 for red cells because transmitted infections still 10 10 questions this morning, you were talking about occur. As I've said, we had hepatitis a few years ago heat treatment and you said "we" -- I think talking 11 11 and we are still at risk of malaria, which can be 12 about haemophilia doctors generally -- "we were crying 12 fatal, and other organisms, bacterial organisms such 13 13 out for pathogen reduction techniques, talking about as Serratia, staph aureus, et cetera, which can also 14 it for years. We wanted it. We needed it". 14 15 Can you tell me a little bit more about the 15 So the answer to your question is that in the 16 years during which you were calling out for pathogen 16 years prior to 1983/84 we knew that hepatitis B was 17 17 reduction techniques and what you had in mind a problem and that it wasn't going away. It still 18 18 particularly. hasn't gone away and we needed something that was 19 A. You may be very surprised to know that that's exactly 19 safer with that. We did not foresee, I don't think 20 what we're discussing at this very moment -- not 20 anybody did really, to be honest, HIV coming down the 21 today, tomorrow probably. Pathogen reduction 21 line. It was out of a clear blue sky, as far as I'm 22 22 techniques for -- transfusion transmitted infections concerned anyway. 23 are still a major problem. They are going to continue 23 As soon as that arrived, it just kick-started 24 to be a major problem and they are very unlikely to go 24 things at long last. Prior to that we had been 25 away and Zika virus -- it usually takes a crisis for 25 discussing hep B and then non-A, non-B hepatitis. 149 150 1 From 1980 onwards, certainly at the Royal Free, 1 that non-A, non-B hepatitis was known about. The peak 2 2 I remember speaking to Dr Kernoff and him saying we incidence was around about 1968, in fact. You didn't 3 need some sort of -- is it going to be some detergent, 3 see it much in children because for some reason they 4 4 heat treatment? What is it going to be? Ultraviolet very rarely have acute episodes but it had been known 5 light? There were lots of discussions at the time but 5 about the adults for a long time and we didn't know 6 there seemed to be no imperative until the crisis and 6 whether it was going to be a very minor thing, 7 7 although that was often the consensus, or whether we had wanted it for years. 8 8 We had wanted self-sufficiency and we had after 1980 it was going to be a prevalent serious 9 9 wanted pathogen reduction techniques, and we had thing or a rare serious thing. But whatever, we 10 wanted pathogen reduction techniques that did not 10 needed -- there were a lot of people, including in the 11 compromise the supply and did not cause inhibitors. 11 haemophilia fraternity, who said, you know, "We need 12 That was part of the problem because all pathogen 12 to be careful about the next thing that comes down the 13 reduction techniques do affect a little, thankfully, 13 line". I don't know that they were expecting anything although initially we thought it was much more, do 14 like HIV, but they were certainly thinking about it. 14 15 15 There's no doubt about that. affect the supply and the supply was crucial and 16 16 I certainly didn't want to go back to using commercial So there was a clamour, yes. But we 17 concentrates again. 17 couldn't -- we couldn't even get to self-sufficiency, 18 I hope that answers it to an extent. 18 let alone reducing further the amount that we had. 19 SIR BRIAN LANGSTAFF: To an extent, yes, you have 19 Sorry. 20 described I think a clamour, certainly from 1980 20 SIR BRIAN LANGSTAFF: So is it the case that, more or 21 onwards, but the implication of years and going back 21 less, for as long as you had been practising in 22 22 to the arguments about self-sufficiency might suggest medicine, in training first and then later on 23 23 it was earlier still. practising, there had been some discussion about

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pathogen reduction techniques?

A. Oh, yes, absolutely, including -- well, because at

A. Oh, yes, it was. Well, because we did know about --

as you know, you've been through it in great detail,

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1 least in part because of the hepatitis B problem. A. It was a probability but again I am sorry to keep 2 2 SIR BRIAN LANGSTAFF: But your sense was until 1980 anyway harking back to prevalence and so on, we didn't know 3 3 what proportion. It ended up something like 16 -- it little imperative to get on with it? 4 A. I think people like Dr Kernoff and so on were very 4 was less in Scotland than in England. It was 5 keen for it to happen. I remember him saying to me, 5 16 per cent or was it 20 per cent or something. But 6 6 you know, the first dose of this concentrate's going the fact is that yes, it was very likely by that stage 7 to give people non-A, non-B hepatitis. It was really 7 in my estimation that it was a transfusion-transmitted 8 8 a question of whether this was going to be like infection. We did not know in haemophilia as yet how 9 9 hepatitis E or G or D or A where it was something that serious this was going to be and how -- what 10 was just eradicated in everyone or nearly everyone or 10 proportion would develop AIDS and how long that would 11 it was going to be thankfully not 100 per cent but 11 take or whatever, or whether it was going to be 12 60 per cent already whatever it was who developed 12 eradicable in some people, like hepatitis C is. That 13 13 turned out to be a false hope unfortunately. chronic problems. 14 It was always regarded, in the people that 14 SIR BRIAN LANGSTAFF: The next question is in relation to 15 15 I spoke to, was always regarded as important. something you were quite self-critical about and that 16 16 SIR BRIAN LANGSTAFF: Thank you. You told us that your is where you told us that you should have done it, 17 view in May 1983, this is I think by reference to what 17 that is pre-test counselling, better. 18 18 Montagnier had been saying in the Pasteur Institute, In what way would you have done pre-test 19 that it was more and more likely -- that was the 19 counselling better, albeit looking at it in 20 expression you used -- that HIV, you said, I think LAV 20 retrospect? 21 is what he called it, had been transmitted and you 21 A. Really, I can deal with it well in retrospect, 22 22 corrected the tense, but the "more and more likely", I think, because it was in the forefront of my mind 23 does that mean you thought it probable or it was just 23 when hepatitis C came along in '89/'90: how do we do 24 a possibility which was becoming a greater possible or 24 this and how do I not repeat history and not learn

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what?

difficult because we didn't have a test that was reliable up until we got the results. So what I should have done better was to contact virology to say: have you got samples? Will you test them? Will you let me know? I understand you're testing various things for Tedder and people to see if it's specific and sensitive and so on.

Then go back to the families and say, "Right, we are going to test this sample as soon as we know, and Dr Tedder is going to let me know when that is likely to be". And then that opens up a whole series of questions which were not adequately dealt with at the time, which are, you know: what if it's positive? What if -- you know, if it's negative, is that fine? Is it negative forever? Can you be infected even now? You know, all those sort of questions would have been much better dealt with with pre-test counselling, and I greatly regret that I didn't spend more time -- or at least realised that there was a gap at the time.

doing it in that optimal way may have been?

A. Well, I think it would be probably have come as less of a shock to the families because they would have been better informed about the potential outcomes. I've said throughout, and it really is a difficult

SIR BRIAN LANGSTAFF: What do you think the effects of not

from it? Basically, what we should have -- it was 154

situation, telling people about potential terrible news when actually it might well not be relevant. It's not an easy thing to do, to deal with uncertainty which goes from one pole to another: life-changing to just carry on. It would, I believe, have -- there were no complaints at the time and I'm amazed. I suppose they were just so dependent upon us that they would not complain. And that's a matter of even greater regret. But the fact is that, you know, they would have been better informed, they would have been stimulated, if you like, to ask more questions and to be better informed. Although there were lots of other means of information, I think it would have been a much better situation, and it's not something that I would have -- I'd feel in any way comfortable about ethically retrospectively. Despite the fact that I can't remember ever asking for this to be done, it was needed.

SIR BRIAN LANGSTAFF: Finally, you have a perspective which few of the witnesses to this Inquiry have in two respects. One is having been so involved during your working life in the Blood Transfusion Service as well as in paediatric haematology and the care of children with haemophilia, but also having done it in Ireland as well as in this country.

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1	A. Yes.	1	anything from it from the local profession point of
1 2	SIR BRIAN LANGSTAFF: By Ireland, I mean southern or the	1 2	anything from it from the legal profession point of view, because of course now it has become virtually
3	Republic of Ireland.	3	impossible, through massive cost and through the
4	A. Yes, since 1940s.	4	Supreme Court judgments here, that you can't really
5	SIR BRIAN LANGSTAFF: Yes.	5	name and shame people anywhere near as it may be
6	A. Still a member of the European Union.	6	necessary in inquiries here. But a lot of good came
7	SIR BRIAN LANGSTAFF: Yes, well, my grandfather was an	7	out of the Lindsay Tribunal, the most important of
8	Irishman, so I know parts of it quite well.	8	which were the resources and the governance and
9	Are there any lessons that we can learn here	9	quality control.
10	from your experiences, what your experiences tell you,	10	I mean, you might say I'm still working in
11	about the way in which Irish Republic has dealt with	11	blood transfusion that blood transfusion has become
12	matters or is dealing with matters?	12	extremely risk-averse, to a level that at times makes
13	A. You may not like the answer to this. I think that the	13	me worry about efficacy and so on and so forth. In
14	fact that the Lindsay Tribunal was held at a time when	14	fact, the blood supply in Ireland now, because of
15	people's memories were better and where data	15	things that are good that have come out of the
16	collection and it was fresh in many people's minds.	16	previous inquiries, and I hope this Inquiry too, is
17	I mean, it's still, to a certain extent, fresh, but	17	that the blood supply now is often on a tightrope.
18	it's a lot of what I say to you, and I have to be	18	And we've just been through an extremely taxing
19	perfectly honest, is relearned memory. And I really	19	period, for myself and others, where the supply has
20	hate that, because what I want to tell you, and I mean	20	been extremely compromised.
21	this, is my actual memory at the time, rather than	21	The Health Life questionnaires now mean that
22	reading through hundreds and hundreds of pages that	22	we which are applied very, very rigorously and
23	I get sent.	23	looked at externally and legally, you know, it's
24	The fact is that the Lindsay Tribunal was	24	100 per cent traceability is a legal requirement in
25	extremely well carried out. You don't want to learn	25	Ireland now. I'm not sure that it is in England. You
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1	have to be sorry, I'm rambling slightly, but in the	1	it is very, very tight indeed, and the management of
2	Ognall Inquiry, that I dealt with, one of the biggest	2	such patients is, as far as I'm concerned, optimal and
3	problems was things like records from Great Ormond	3	a lot of that came because of that inquiry. A lot of
4	Street, where are they? Who had what? How can you do	4	good came of it.
5	a look-back when you don't know what people were	5	SIR BRIAN LANGSTAFF: Those are all the questions that
6	seeing, et cetera?	6	I have to ask.
7	So of that came the product liability or what	7	MS RICHARDS: Professor Hann, is there anything further
8	I call product liability, Consumer Protection Act,	8	that you wanted to add?
9	et cetera, within Ireland now. The similar thing	9	A. Well, I went to say two things really. First of all,
10	happened because of Lindsay and so on. And we now	10	and I will be very brief, I want to thank you and
11	have an absolute requirement for everyone who uses	11	Sir Brian for letting me ramble on in a very verbose
12	blood products to have complete traceability, which	12	way sometimes because I have two pages of things I was
13	has to be verified and annotated.	13	going to say here and I have already, I think, said
14	Sorry, the final thing, really, is that they	14	them all.
15	went from having one haemophilia doctor to having	15	I do think it's very important that the Inquiry
16	a whole cohort now and a proper built centre and	16	takes on board the special requirements for children
17	excellent governance. It's revolutionised haemophilia	17	and paediatric care and paediatric haemophilia care
18	care here. I think it is there's no doubt that you	18	and all those aspects that I've talked about, and
19	will find, I'm sure, it difficult to make judgments of	19	I too hope that good comes of this for reasons I've
20	things that happened a very long time ago. They were	20	already stated. It's an ongoing problem and an
21	in a better position in that respect and it's possible	21	existential problem.
22	that the record-keeping itself was better here than	22	The final thing I want to say is you have
23	there.	23	already heard a lot of people say that the worst thing
24	But from the Irish point of view, haemophilia	24	you can do, if you are in your right mind as a doctor,
25	care now is extremely well run. The governance over	25	is to do harm to people. Unfortunately, harm does

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1	occur. A tragedy such as this is not absolutely	1	evidence restless for improvement. So I'd like to
2	unique but it was unique in my experience and it was	2	thank you in particular for all that and I'd ask you
3	a terrible thing and it caused terrible suffering to	3	for your part to forgive me and us for having taken up
4	the families, and I just want to apologise again for	4	your day and prevented you from getting out to enjoy
5	the additional harm that I caused through not	5	what may be a rare Irish experience, that of sunshine
6	communicating well enough. Thank you.	6	as you told us this morning.
7	MS RICHARDS: Thank you, professor.	7	A. Thank you. I am very grateful for the opportunity.
8	SIR BRIAN LANGSTAFF: Professor, you have given us a very	8	MS RICHARDS: Sir, we resume tomorrow at 2.00 with the
9	different perspective in many ways from the	9	commencement of the evidence of Professor Lowe.
10	perspectives we've been listening to because yours is	10	SIR BRIAN LANGSTAFF: So 2 o'clock tomorrow afternoon
11	the perspective of somebody who is a paediatric	11	(4.19 pm)
12	haemophilia doctor in part, if I can call you that.	12	(Adjourned until 2.00 pm the following day)
13	You've not held back at all in what you have	13	
14	had to say and your views, and it seemed to me that	14	
15	you have given us a very graphic picture of what it	15	
16	was like to start in 1983 with a huge range of	16	
17	responsibilities and very little resource and a system	17	
18	which needed to be changed as you saw it.	18	
19	You have I think shown that your approach was	19	
20	to do it, not as being the King who is a doctor or	20	
21	a consultant but as part of the skilled team working	21	
22	together, which you described right at the end of your	22	
23	evidence as the ethos which you wanted to instil and	23	
24	it struck me that you have been restless throughout	24	
25	your career as you've told us and, indeed, in your	25	
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